

PCTWORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau

INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶ : C07C 235/84, C07D 213/55, 239/28, 241/24, A61K 31/17, 31/44, 31/50, 31/505		A1	(11) International Publication Number: WO 99/46236
			(43) International Publication Date: 16 September 1999 (16.09.99)
(21) International Application Number: PCT/DK99/00122			(72) Inventors: RICHTER, Lutz Stefan (deceased); ANDERSEN, Henrik, Sune; Gustav Adolphsvej 2, DK-2800 Lyngby (DK). VAGNER, Josef; Jonstrupvej 271A, DK-3500 Vaerloese (DK). JEPPESEN, Claus, Bekker; Damgaardsvej 17, Niverød, DK-2990 Nivaa (DK). MØLLER, Niels, Peter, Hundahl; Midtermolen 4, 3, DK-2100 Copenhagen Ø (DK). BRANNER, Sven; Ved Smedebakken 7A, DK-2800 Lyngby (DK). SU, Jing; Apartment 4A, 220 E. 70 Street, New York, NY 10021 (US). BAKIR, Farid; Apartment A, 5757 College Avenue, San Diego, CA 92120 (US). JUDGE, Luke, Milburn; 644 1/2 Bonair Way, La Jolla, CA 92037 (US).
(22) International Filing Date: 11 March 1999 (11.03.99)			
(30) Priority Data: 0342/98 12 March 1998 (12.03.98) DK 0345/98 12 March 1998 (12.03.98) DK 0472/98 3 April 1998 (03.04.98) DK 0479/98 3 April 1998 (03.04.98) DK PA 1998 00940 15 July 1998 (15.07.98) DK			
(71) Applicants: NOVO NORDISK A/S [DK/DK]; Corporate Patents, Novo Allé, DK-2880 Bagsvaerd (DK). ONTOGEN CORPORATION [US/US]; 6451 El Camino Real, Carlsbad, CA 92009 (US).			
(74) Common Representative: NOVO NORDISK A/S; Corporate Patents, Novo Allé, DK-2880 Bagsvaerd (DK).			
(81) Designated States: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).			
Published <i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>			
(54) Title: MODULATORS OF PROTEIN TYROSINE PHOSPHATASES (PTPASES)			
(57) Abstract <p>The present invention provides novel compounds, novel compositions, methods of their use, and methods of their manufacture, where such compounds are pharmacologically useful inhibitors of Protein Tyrosine Phosphatases (PTPases) such as PTP1B, CD45, SHP-1, SHP-2, PTPα, LAR and HePTP or the like. The compounds are useful in the treatment of type I diabetes, type II diabetes, impaired glucose tolerance, insulin resistance, obesity, immune dysfunctions including autoimmunity diseases with dysfunctions of the coagulation system, allergic diseases including asthma, osteoporosis, proliferative disorders including cancer and psoriasis, diseases with decreased or increased synthesis or effects of growth hormone, diseases with decreased or increased synthesis of hormones or cytokines that regulate the release of/or response to growth hormone, diseases of the brain including Alzheimer's disease and schizophrenia, and infectious diseases.</p>			

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

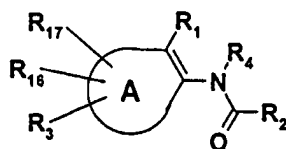
AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakstan	RO	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

Modulators of Protein Tyrosine Phosphatases (PTPases)

FIELD OF THE INVENTION

- 5 The present invention relates to novel compounds, to methods for their preparation, to compositions comprising the compounds, to the use of these compounds as medicaments and their use in therapy, where such compounds of Formula 1 are pharmacologically useful inhibitors of Protein Tyrosine Phosphatases (PTPases) such as PTP1B, CD45, SHP-1, SHP-2, PTP α , LAR and HePTP or the like,

10



Formula 1

wherein A, R₁, R₂, R₃, R₄, R₁₆ and R₁₇ are defined more fully below.

- 15 It has been found that PTPases plays a major role in the intracellular modulation and regulation of fundamental cellular signalling mechanisms involved in metabolism, growth, proliferation and differentiation (Flint et al., The EMBO J. 12:1937-46 (1993); Fischer et al, Science 253:401-6 (1991)). Overexpression or altered activity of tyrosine phosphatases can also contribute to the symptoms and progression of various
- 20 diseases (Wiener, et al., J. Natl. cancer Inst. 86:372-8 (1994); Hunter and Cooper, Ann. Rev. Biochem, 54:897-930 (1985)). Furthermore, there is increasing evidence which suggests that inhibition of these PTPases may help treat certain types of diseases such as diabetes type I and II, autoimmune disease, acute and chronic inflammation, osteoporosis and various forms of cancer.

25

BACKGROUND OF THE INVENTION

Protein phosphorylation is now well recognized as an important mechanism utilized
5 by cells to transduce signals during different stages of cellular function (Fischer et al, Science 253:401-6 (1991); Flint et al., The EMBO J. 12:1937-46 (1993)). There are at least two major classes of phosphatases: (1) those that dephosphorylate proteins (or peptides) that contain a phosphate group(s) on a serine or threonine moiety (termed Ser/Thr phosphatases) and (2) those that remove a phosphate group(s)
10 from the amino acid tyrosine (termed protein tyrosine phosphatases or PTPases).

The PTPases are a family of enzymes that can be classified into two groups: a) intracellular or nontransmembrane PTPases and b) receptor-type or transmembrane PTPases.

15

Intracellular PTPases: Most known intracellular type PTPases contain a single conserved catalytic phosphatase domain consisting of 220-240 amino acid residues. The regions outside the PTPase domains are believed to play important roles in localizing the intracellular PTPases subcellularly (Mauro, L.J. and Dixon, J.E. *TIBS* 19: 151-155 (1994)). The first intracellular PTPase to be purified and
20 characterized was **PTP1B** which was isolated from human placenta (Tonks et al., *J. Biol. Chem.* 263: 6722-6730 (1988)). Shortly after, PTP1B was cloned (Charbonneau et al., *Proc. Natl. Acad. Sci. USA* 86: 5252-5256 (1989); Chernoff et al., *Proc. Natl. Acad. Sci. USA* 87: 2735-2789 (1989)). Other examples of
25 intracellular PTPases include (1) **T-cell PTPase** (Cool et al. *Proc. Natl. Acad. Sci. USA* 86: 5257-5261 (1989)), (2) **rat brain PTPase** (Guan et al., *Proc. Natl. Acad. Sci. USA* 87:1501-1502 (1990)), (3) neuronal phosphatase **STEP** (Lombroso et al., *Proc. Natl. Acad. Sci. USA* 88: 7242-7246 (1991)), (4) ezrin-domain containing PTPases: **PTPMEG1** (Guet et al., *Proc. Natl. Acad. Sci. USA* 88: 5867-
30 57871 (1991)), **PTPH1** (Yang and Tonks, *Proc. Natl. Acad. Sci. USA* 88: 5949-5953 (1991)), **PTPD1** and **PTPD2** (Møller et al., *Proc. Natl. Acad. Sci. USA* 91:

7477-7481 (1994)), **FAP-1/BAS** (Sato *et al.*, *Science* 268: 411-415 (1995); Banville *et al.*, *J. Biol. Chem.* 269: 22320-22327 (1994); Maekawa *et al.*, *FEBS Letters* 337: 200-206 (1994)), and SH2 domain containing PTPases: **PTP1C/SH-PTP1/SHP-1** (Plutzky *et al.*, *Proc. Natl. Acad. Sci. USA* 89: 1123-1127 (1992);
5 Shen *et al.*, *Nature Lond.* 352: 736-739 (1991)) and **PTP1D/Syp/SH-PTP2/SHP-2** (Vogel *et al.*, *Science* 259: 1611-1614 (1993); Feng *et al.*, *Science* 259: 1607-1611 (1993); Bastein *et al.*, *Biochem. Biophys. Res. Comm.* 196: 124-133 (1993)).

Low molecular weight phosphotyrosine-protein phosphatase (**LMW-PTPase**)
10 shows very little sequence identity to the intracellular PTPases described above. However, this enzyme belongs to the PTPase family due to the following characteristics: (i) it possesses the PTPase active site motif: Cys-Xxx-Xxx-Xxx-Xxx-Xxx-Arg (Cirri *et al.*, *Eur. J. Biochem.* 214: 647-657 (1993)); (ii) this Cys
residue forms a phospho-intermediate during the catalytic reaction similar to the
15 situation with 'classical' PTPases (Cirri *et al.*, *supra*; Chiarugi *et al.*, *FEBS Lett.* 310: 9-12 (1992)); (iii) the overall folding of the molecule shows a surprising degree of similarity to that of PTP1B and *Yersinia* PTP (Su *et al.*, *Nature* 370: 575-578 (1994)).

20 *Receptor-type PTPases* consist of a) a putative ligand-binding extracellular domain, b) a transmembrane segment, and c) an intracellular catalytic region. The structures and sizes of the putative ligand-binding extracellular domains of receptor-type PTPases are quite divergent. In contrast, the intracellular catalytic regions of receptor-type PTPases are very homologous to each other and to the
25 intracellular PTPases. Most receptor-type PTPases have two tandemly duplicated catalytic PTPase domains.

The first receptor-type PTPases to be identified were (1) **CD45/LCA** (Ralph, S.J., *EMBO J.* 6: 1251-1257 (1987)) and (2) **LAR** (Streuli *et al.*, *J. Exp. Med.* 168:
30 1523-1530 (1988)) that were recognized to belong to this class of enzymes based on homology to PTP1B (Charbonneau *et al.*, *Proc. Natl. Acad. Sci. USA* 86: 5252-

5256 (1989)). CD45 is a family of high molecular weight glycoproteins and is one of the most abundant leukocyte cell surface glycoproteins and appears to be exclusively expressed upon cells of the hematopoietic system (Trowbridge and Thomas, *Ann. Rev. Immunol.* 12: 85-116 (1994)).

5

The identification of CD45 and LAR as members of the PTPase family was quickly followed by identification and cloning of several different members of the receptor-type PTPase group. Thus, 5 different PTPases, (3) PTP α , (4) PTP β , (5) PTP δ , (6) PTP ϵ , and (7) PTP ζ , were identified in one early study (Krueger *et al.*, *EMBO J.* 9: 3241-3252 (1990)). Other examples of receptor-type PTPases include (8) PTP γ (Barnea *et al.*, *Mol. Cell. Biol.* 13: 1497-1506 (1995)) which, like PTP ζ (Krueger and Saito, *Proc. Natl. Acad. Sci. USA* 89: 7417-7421 (1992)) contains a carbonic anhydrase-like domain in the extracellular region, (9) PTP μ (Gebbink *et al.*, *FEBS Letters* 290: 123-130 (1991)), (10) PTP κ (Jiang *et al.*, *Mol. Cell. Biol.* 13: 2942-2951 (1993)). Based on structural differences the receptor-type PTPases may be classified into subtypes (Fischer *et al.*, *Science* 253: 401-406 (1991)): (I) CD45; (II) LAR, PTP δ , (11) PTP σ ; (III) PTP β , (12) SAP-1 (Matozaki *et al.*, *J. Biol. Chem.* 269: 2075-2081 (1994)), (13) PTP-U2/GLEPP1 (Seimiya *et al.*, *Oncogene* 10: 1731-1738 (1995); Thomas *et al.*, *J. Biol. Chem.* 269: 19953-19962 (1994)), and (14) DEP-1; (IV) PTP α , PTP ϵ . All receptor-type PTPases except Type IV contain two PTPase domains. Novel PTPases are continuously identified, and it is anticipated that more than 500 different species will be found in the human genome, i.e. close to the predicted size of the protein tyrosine kinase superfamily (Hanks and Hunter, *FASEB J.* 9: 576-596 (1995)).

25

PTPases are the biological counterparts to protein tyrosine kinases (PTKs). Therefore, one important function of PTPases is to control, down-regulate, the activity of PTKs. However, a more complex picture of the function of PTPases now emerges. Several studies have shown that some PTPases may actually act as positive mediators of cellular signalling. As an example, the SH2 domain-containing PTP1D seems to act as a positive mediator in insulin-stimulated Ras

30

activation (Noguchi *et al.*, *Mol. Cell. Biol.* 14: 6674-6682 (1994)) and of growth factor-induced mitogenic signal transduction (Xiao *et al.*, *J. Biol. Chem.* 269: 21244-21248 (1994)), whereas the homologous PTP1C seems to act as a negative regulator of growth factor-stimulated proliferation (Bignon and
5 Siminovitch, *Clin. Immunol. Immunopathol.* 73: 168-179 (1994)). Another example of PTPases as positive regulators has been provided by studies designed to define the activation of the Src-family of tyrosine kinases. In particular, several lines of evidence indicate that CD45 is positively regulating the activation of hematopoietic cells, possibly through dephosphorylation of the C-terminal tyrosine
10 of Fyn and Lck (Chan *et al.*, *Annu. Rev. Immunol.* 12: 555-592 (1994)).

Dual specificity protein tyrosine phosphatases (dsPTPases) define a subclass within the PTPases family that can hydrolyze phosphate from phosphotyrosine as well as from phosphor-serine/threonine. dsPTPases contain the signature
15 sequence of PTPases: His-Cys-Xxx-Xxx-Gly-Xxx-Xxx-Arg. At least three dsPTPases have been shown to dephosphorylate and inactivate extracellular signal-regulated kinase (ERKs)/mitogen-activated protein kinase (MAPK): **MAPK phosphatase** (CL100, 3CH134) (Charles *et al.*, *Proc. Natl. Acad. Sci. USA* 90: 5292-5296 (1993)); **PAC-1** (Ward *et al.*, *Nature* 367: 651-654 (1994)); **rVH6**
20 (Mourey *et al.*, *J. Biol. Chem.* 271: 3795-3802 (1996)). Transcription of dsPTPases are induced by different stimuli, e.g. oxidative stress or heat shock (Ishibashi *et al.*, *J. Biol. Chem.* 269: 29897-29902 (1994); Keyse and Emslie, *Nature* 359: 644-647 (1992)). Further, they may be involved in regulation of the cell cycle: **cdc25** (Millar and Russell, *Cell* 68: 407-410 (1992)); **KAP** (Hannon *et al.*, *Proc. Natl. Acad. Sci. USA* 91: 1731-1735 (1994)). Interestingly, tyrosine dephosphorylation of cdc2 by a
25 dual specific phosphatase, cdc25, is required for induction of mitosis in yeast (review by Walton and Dixon, *Annu. Rev. Biochem.* 62: 101-120 (1993)).

PTPases were originally identified and purified from cell and tissue lysates using a
30 variety of artificial substrates and therefore their natural function of dephosphorylation was not well known. Since tyrosine phosphorylation by tyrosine kinases is usually

associated with cell proliferation, cell transformation and cell differentiation, it was assumed that PTPases were also associated with these events.

This association has now been proven to be the case with many PTPases. PTP1B, a phosphatase whose structure was recently elucidated (Barford et al., Science 263:1397-1404 (1994)) has been shown to be involved in insulin-induced oocyte maturation (Flint et al., The EMBO J. 12:1937-46 (1993)) and recently it has been suggested that the overexpression of this enzyme may be involved in p185^{c-erb B2} - associated breast and ovarian cancers (Wiener, et al., J. Natl. cancer Inst. 86:372-8 (1994); Weiner et al., Am. J. Obstet. Gynecol. 170:1177-883 (1994)). The insulin-induced oocyte maturation mechanism has been correlated with the ability of PTP1B to block activation of S6 kinase. The association with cancer is recent evidence which suggests that overexpression of PTP1B is statistically correlated with increased levels of p185^{c-erb B2} in ovarian and breast cancer. The role of PTP1B in the etiology and progression of the disease has not yet been elucidated. Inhibitors of PTP1B may therefore help clarify the role of PTP1B in cancer and in some cases provide therapeutic treatment for certain forms of cancer.

The activity of a number of other newly discussed phosphatases are currently under investigation. Two of these: SHP-1 and Syp/PTP1D/SHPTP2/PTP2C/SHP-2 have recently been implicated in the activation of Platelet Derived Growth Factor and Epidermal Growth Factor induced responses (Li et al., Mole. Cell. Biol. 14:509-17 (1994)). Since both growth factors are involved in normal cell processing as well as disease states such as cancer and arteriosclerosis, it is hypothesized that inhibitors of these phosphatases would also show therapeutic efficacy. Accordingly, the compounds of the present invention which exhibit inhibitory activity against various PTPases, are indicated in the treatment or management of the foregoing diseases.

PTPases: the insulin receptor signalling pathway/diabetes

Insulin is an important regulator of different metabolic processes and plays a key role in the control of blood glucose. Defects related to its synthesis or signalling

lead to diabetes mellitus. Binding of insulin to its receptor causes rapid (auto)phosphorylation of several tyrosine residues in the intracellular part of the b-subunit. Three closely positioned tyrosine residues (the tyrosine-1150 domain) must all be phosphorylated to obtain full activity of the insulin receptor tyrosine kinase (IRTK) which transmits the signal further downstream by tyrosine phosphorylation of other cellular substrates, including insulin receptor substrate-1 (IRS-1) (Wilden *et al.*, *J. Biol. Chem.* 267: 16660-16668 (1992); Myers and White, *Diabetes* 42: 643-650 (1993); Lee and Pilch, *Am. J. Physiol.* 266: C319-C334 (1994); White *et al.*, *J. Biol. Chem.* 263: 2969-2980 (1988)). The structural basis for the function of the tyrosine-triplet has been provided by recent X-ray crystallographic studies of IRTK that showed tyrosine-1150 to be autoinhibitory in its unphosphorylated state (Hubbard *et al.*, *Nature* 372: 746-754 (1994)).

Several studies clearly indicate that the activity of the auto-phosphorylated IRTK can be reversed by dephosphorylation *in vitro* (reviewed in Goldstein, *Receptor* 3: 1-15 (1993); Mooney and Anderson, *J. Biol. Chem.* 264: 6850-6857 (1989)), with the tri-phosphorylated tyrosine-1150 domain being the most sensitive target for protein-tyrosine phosphatases (PTPases) as compared to the di- and mono-phosphorylated forms (King *et al.*, *Biochem. J.* 275: 413-418 (1991)). It is, therefore, tempting to speculate that this tyrosine-triplet functions as a control switch of IRTK activity. Indeed, the IRTK appears to be tightly regulated by PTP-mediated dephosphorylation *in vivo* (Khan *et al.*, *J. Biol. Chem.* 264: 12931-12940 (1989); Faure *et al.*, *J. Biol. Chem.* 267: 11215-11221 (1992); Rothenberg *et al.*, *J. Biol. Chem.* 266: 8302-8311 (1991)). The intimate coupling of PTPases to the insulin signalling pathway is further evidenced by the finding that insulin differentially regulates PTPase activity in rat hepatoma cells (Meyerovitch *et al.*, *Biochemistry* 31: 10338-10344 (1992)) and in livers from alloxan diabetic rats (Boylan *et al.*, *J. Clin. Invest.* 90: 174-179 (1992)).

Relatively little is known about the identity of the PTPases involved in IRTK regulation. However, the existence of PTPases with activity towards the insulin

receptor can be demonstrated as indicated above. Further, when the strong PTPase-inhibitor pervanadate is added to whole cells an almost full insulin response can be obtained in adipocytes (Fantus *et al.*, *Biochemistry* 28: 8864-8871 (1989); Eriksson *et al.*, *Diabetologia* 39: 235-242 (1995)) and skeletal muscle (Leighton *et al.*, *Biochem. J.* 276: 289-292 (1991)). In addition, recent studies show that a new class of peroxovanadium compounds act as potent hypoglycemic compounds *in vivo* (Posner *et al.*, *supra*). Two of these compounds were demonstrated to be more potent inhibitors of dephosphorylation of the insulin receptor than of the EGF-receptor.

It was recently found that the ubiquitously expressed SH2 domain containing PTPase, PTP1D (Vogel *et al.*, 1993, *supra*), associates with and dephosphorylates IRS-1, but apparently not the IR itself (Kuhné *et al.*, *J. Biol. Chem.* 268: 11479-11481 (1993); (Kuhné *et al.*, *J. Biol. Chem.* 269: 15833-15837 (1994)).

Previous studies suggest that the PTPases responsible for IRTK regulation belong to the class of membrane-associated (Faure *et al.*, *J. Biol. Chem.* 267: 11215-11221 (1992)) and glycosylated molecules (Häring *et al.*, *Biochemistry* 23: 3298-3306 (1984); Sale, *Adv. Prot. Phosphatases* 6: 159-186 (1991)). Hashimoto *et al.* have proposed that LAR might play a role in the physiological regulation of insulin receptors in intact cells (Hashimoto *et al.*, *J. Biol. Chem.* 267: 13811-13814 (1992)). Their conclusion was reached by comparing the rate of dephosphorylation/inactivation of purified IR using recombinant PTP1B as well as the cytoplasmic domains of LAR and PTPa. Antisense inhibition was recently used to study the effect of LAR on insulin signalling in a rat hepatoma cell line (Kulas *et al.*, *J. Biol. Chem.* 270: 2435-2438 (1995)). A suppression of LAR protein levels by about 60 percent was paralleled by an approximately 150 percent increase in insulin-induced auto-phosphorylation. However, only a modest 35 percent increase in IRTK activity was observed, whereas the insulin-dependent phosphatidylinositol 3-kinase (PI 3-kinase) activity was significantly increased by 350 percent. Reduced LAR levels did not alter the basal level of IRTK tyrosine

phosphorylation or activity. The authors speculate that LAR could specifically dephosphorylate tyrosine residues that are critical for PI 3-kinase activation either on the insulin receptor itself or on a downstream substrate.

- 5 While previous reports indicate a role of PTPa in signal transduction through src activation (Zheng *et al.*, *Nature* 359: 336-339 (1992); den Hertog *et al.*, *EMBO J.* 12: 3789-3798 (1993)) and interaction with GRB-2 (den Hertog *et al.*, *EMBO J.* 13: 3020-3032 (1994); Su *et al.*, *J. Biol. Chem.* 269: 18731-18734 (1994)), a recent study suggests a function for this phosphatase and its close relative PTPe as
- 10 negative regulators of the insulin receptor signal (Møller *et al.*, 1995 *supra*). This study also indicates that receptor-like PTPases play a significant role in regulating the IRTK, whereas intracellular PTPases seem to have little, if any, activity towards the insulin receptor. While it appears that the target of the negative regulatory activity of PTPases a and e is the receptor itself, the downmodulating
- 15 effect of the intracellular TC-PTP seems to be due to a downstream function in the IR-activated signal. Although PTP1B and TC-PTP are closely related, PTP1B had only little influence on the phosphorylation pattern of insulin-treated cells. Both PTPases have distinct structural features that determine their subcellular localization and thereby their access to defined cellular substrates (Frangione *et*
- 20 *al.*, *Cell* 68: 545-560 (1992); Faure and Posner, *Glia* 9: 311-314 (1993)). Therefore, the lack of activity of PTP1B and TC-PTP towards the IRTK may, at least in part, be explained by the fact that they do not co-localize with the activated insulin receptor. In support of this view, PTP1B and TC-PTP have been excluded as candidates for the IR-associated PTPases in hepatocytes based on subcellular
- 25 localization studies (Faure *et al.*, *J. Biol. Chem.* 267: 11215-11221 (1992)).

The transmembrane PTPase CD45, which is believed to be hematopoietic cell-specific, was in a recent study found to negatively regulate the insulin receptor tyrosine kinase in the human multiple myeloma cell line U266 (Kulas *et al.*, *J. Biol.*

30 *Chem.* 271: 755-760 (1996)).

PTPases: somatostatin

Somatostatin inhibits several biological functions including cellular proliferation (Lamberts *et al.*, *Molec. Endocrinol.* 8: 1289-1297 (1994)). While part of the antiproliferative activities of somatostatin are secondary to its inhibition of hormone and growth factor secretion (e.g. growth hormone and epidermal growth factor), other antiproliferative effects of somatostatin are due to a direct effect on the target cells. As an example, somatostatin analogs inhibit the growth of pancreatic cancer presumably via stimulation of a single PTPase, or a subset of PTPases, rather than a general activation of PTPase levels in the cells (Liebow *et al.*, *Proc. Natl. Acad. Sci. USA* 86: 2003-2007 (1989); Colas *et al.*, *Eur. J. Biochem.* 207: 1017-1024 (1992)). In a recent study it was found that somatostatin stimulation of somatostatin receptors SSTR1, but not SSTR2, stably expressed in CHO-K1 cells can stimulate PTPase activity and that this stimulation is pertussis toxin-sensitive. Whether the inhibitory effect of somatostatin on hormone and growth factor secretion is caused by a similar stimulation of PTPase activity in hormone producing cells remains to be determined.

PTPases: the immune system/autoimmunity

Several studies suggest that the receptor-type PTPase CD45 plays a critical role not only for initiation of T cell activation, but also for maintaining the T cell receptor-mediated signalling cascade. These studies are reviewed in: (Weiss A., *Ann. Rev. Genet.* 25: 487-510 (1991); Chan *et al.*, *Annu. Rev. Immunol.* 12: 555-592 (1994); Trowbridge and Thomas, *Annu. Rev. Immunol.* 12: 85-116 (1994)). CD45 is one of the most abundant of the cell surface glycoproteins and is expressed exclusively on hemopoietic cells. In T cells, it has been shown that CD45 is one of the critical components of the signal transduction machinery of lymphocytes. In particular, evidence has suggested that CD45 phosphatase plays a pivotal role in antigen-stimulated proliferation of T lymphocytes after an antigen has bound to the T cell receptor (Trowbridge, *Ann. Rev. Immunol.* 12:85-116 (1994)). Several studies suggest

that the PTPase activity of CD45 plays a role in the activation of Lck, a lymphocyte-specific member of the Src family protein-tyrosine kinase (Mustelin et al., Proc. Natl. Acad. Sci. USA 86: 6302-6306 (1989); Ostergaard et al., Proc. Natl. Acad. Sci. USA 86: 8959-8963 (1989)). These authors hypothesized that the phosphatase activity of CD45 activates Lck by dephosphorylation of a C-terminal tyrosine residue, which may, in turn, be related to T-cell activation. In a recent study it was found that recombinant p56lck specifically associates with recombinant CD45 cytoplasmic domain protein, but not to the cytoplasmic domain of the related PTPa (Ng et al., *J. Biol. Chem.* 271: 1295-1300 (1996)). The p56lck-CD45 interaction seems to be mediated via a nonconventional SH2 domain interaction not requiring phosphotyrosine. In immature B cells, another member of the Src family protein-tyrosine kinases, Fyn, seems to be a selective substrate for CD45 compared to Lck and Syk (Katagiri et al., *J. Biol. Chem.* 270: 27987-27990 (1995)).

Studies using transgenic mice with a mutation for the CD45-exon6 exhibited lacked mature T cells. These mice did not respond to an antigenic challenge with the typical T cell mediated response (Kishihara et al., *Cell* 74:143-56 (1993)). Inhibitors of CD45 phosphatase would therefore be very effective therapeutic agents in conditions that are associated with autoimmune disease.

CD45 has also been shown to be essential for the antibody mediated degranulation of mast cells (Berger et al., *J. Exp. Med.* 180:471-6 (1994)). These studies were also done with mice that were CD45-deficient. In this case, an IgE-mediated degranulation was demonstrated in wild type but not CD45-deficient T cells from mice. These data suggest that CD45 inhibitors could also play a role in the symptomatic or therapeutic treatment of allergic disorders.

Another recently discovered PTPase, an inducible lymphoid-specific protein tyrosine phosphatase (HePTP) has also been implicated in the immune response. This phosphatase is expressed in both resting T and B lymphocytes, but not non-hemopoietic cells. Upon stimulation of these cells, mRNA levels from the HePTP gene increase 10-15 fold (Zanke et al., *Eur. J. Immunol.* 22:235-239 (1992)). In both T and B cells

HePTP may function during sustained stimulation to modulate the immune response through dephosphorylation of specific residues. Its exact role, however remains to be defined.

5 Likewise, the hematopoietic cell specific PTP1C seems to act as a negative regulator and play an essential role in immune cell development. In accordance with the above-mentioned important function of CD45, HePTP and PTP1C, selective PTPase inhibitors may be attractive drug candidates both as immunosuppressors and as immunostimulants. One recent study illustrates the
10 potential of PTPase inhibitors as immunomodulators by demonstrating the capacity of the vanadium-based PTPase inhibitor, BMLOV, to induce apparent B cell selective apoptosis compared to T cells (Schieven et al., *J. Biol. Chem.* 270: 20824-20831 (1995)).

15 PTPases: cell-cell interactions/cancer

Focal adhesion plaques, an *in vitro* phenomenon in which specific contact points are formed when fibroblasts grow on appropriate substrates, seem to mimic, at least in part, cells and their natural surroundings. Several focal adhesion proteins
20 are phosphorylated on tyrosine residues when fibroblasts adhere to and spread on extracellular matrix (Gumbiner, *Neuron* 11, 551-564 (1993)). However, aberrant tyrosine phosphorylation of these proteins can lead to cellular transformation. The intimate association between PTPases and focal adhesions is supported by the finding of several intracellular PTPases with ezrin-like N-terminal domains, e.g.
25 PTPMEG1 (Gu et al., *Proc. Natl. Acad. Sci. USA* 88: 5867-5871 (1991)), PTPH1 (Yang and Tonks, *Proc. Natl. Acad. Sci. USA* 88: 5949-5953 (1991)) and PTPD1 (Møller et al., *Proc. Natl. Acad. Sci. USA* 91: 7477-7481 (1994)). The ezrin-like domain show similarity to several proteins that are believed to act as links between the cell membrane and the cytoskeleton. PTPD1 was found to be phosphorylated
30 by and associated with c-src *in vitro* and is hypothesized to be involved in the regulation of phosphorylation of focal adhesions (Møller et al., *supra*).

PTPases may oppose the action of tyrosine kinases, including those responsible for phosphorylation of focal adhesion proteins, and may therefore function as natural inhibitors of transformation. TC-PTP, and especially the truncated form of this enzyme (Cool *et al.*, *Proc. Natl. Acad. Sci. USA* 87: 7280-7284 (1990)), can inhibit the transforming activity of v-erb and v-fms (Lammers *et al.*, *J. Biol. Chem.* 268: 22456-22462 (1993); Zander *et al.*, *Oncogene* 8: 1175-1182 (1993)). Moreover, it was found that transformation by the oncogenic form of the *HER2/neu* gene was suppressed in NIH 3T3 fibroblasts overexpressing PTP1B (Brown-Shimer *et al.*, *Cancer Res.* 52: 478-482 (1992)).

The expression level of PTP1B was found to be increased in a mammary cell line transformed with *neu* (Zhay *et al.*, *Cancer Res.* 53: 2272-2278 (1993)). The intimate relationship between tyrosine kinases and PTPases in the development of cancer is further evidenced by the recent finding that PTPe is highly expressed in murine mammary tumors in transgenic mice over-expressing c-*neu* and v-Ha-*ras*, but not c-*myc* or *int-2* (Elson and Leder, *J. Biol. Chem.* 270: 26116-26122 (1995)). Further, the human gene encoding PTPg was mapped to 3p21, a chromosomal region which is frequently deleted in renal and lung carcinomas (LaForgia *et al.*, *Proc. Natl. Acad. Sci. USA* 88: 5036-5040 (1991)).

In this context, it seems significant that PTPases appear to be involved in controlling the growth of fibroblasts. In a recent study it was found that Swiss 3T3 cells harvested at high density contain a membrane-associated PTPase whose activity on an average is 8-fold higher than that of cells harvested at low or medium density (Pallen and Tong, *Proc. Natl. Acad. Sci. USA* 88: 6996-7000 (1991)). It was hypothesized by the authors that density-dependent inhibition of cell growth involves the regulated elevation of the activity of the PTPase(s) in question. In accordance with this view, a novel membrane-bound, receptor-type PTPase, DEP-1, showed enhanced (≥ 10 -fold) expression levels with increasing cell density of WI-38 human embryonic lung fibroblasts and in the AG1518

fibroblast cell line (Östman *et al.*, *Proc. Natl. Acad. Sci. USA* 91: 9680-9684 (1994)).

Two closely related receptor-type PTPases; PTP κ and PTP μ , can mediate
5 homophilic cell-cell interaction when expressed in non-adherent insect cells,
suggesting that these PTPases might have a normal physiological function in cell-
to-cell signalling (Gebbink *et al.*, *J. Biol. Chem.* 268: 16101-16104 (1993); Brady-
Kalnay *et al.*, *J. Cell Biol.* 122: 961-972 (1993); Sap *et al.*, *Mol. Cell. Biol.* 14: 1-9
(1994)). Interestingly, PTP κ and PTP μ do not interact with each other, despite
10 their structural similarity (Zondag *et al.*, *J. Biol. Chem.* 270: 14247-14250 (1995)).
From the studies described above it is apparent that PTPases may play an
important role in regulating normal cell growth. However, as pointed out above,
recent studies indicate that PTPases may also function as positive mediators of
intracellular signalling and thereby induce or enhance mitogenic responses.
15 Increased activity of certain PTPases might therefore result in cellular
transformation and tumor formation. Indeed, in one study over-expression of
PTP α was found to lead to transformation of rat embryo fibroblasts (Zheng, *supra*).
In addition, a novel PTP, SAP-1, was found to be highly expressed in pancreatic
and colorectal cancer cells. SAP-1 is mapped to chromosome 19 region q13.4 and
20 might be related to carcinoembryonic antigen mapped to 19q13.2 (Uchida *et al.*, *J.*
Biol. Chem. 269: 12220-12228 (1994)). Further, the dsPTPase, cdc25,
dephosphorylates cdc2 at Thr14/Tyr-15 and thereby functions as positive regulator
of mitosis (reviewed by Hunter, *Cell* 80: 225-236 (1995)). Inhibitors of specific
PTPases are therefore likely to be of significant therapeutic value in the treatment
25 of certain forms of cancer.

PTPases: platelet aggregation

Recent studies indicate that PTPases are centrally involved in platelet
30 aggregation. Agonist-induced platelet activation results in calpain-catalyzed
cleavage of PTP1B with a concomitant 2-fold stimulation of PTPase activity

(Frangioni *et al.*, *EMBO J.* 12: 4843-4856 (1993)). The cleavage of PTP1B leads to subcellular relocation of the enzyme and correlates with the transition from reversible to irreversible platelet aggregation in platelet-rich plasma. In addition, the SH2 domain containing PTPase, SHP-1, was found to translocate to the cytoskeleton in platelets after thrombin stimulation in an aggregation-dependent manner (Li *et al.*, *FEBS Lett.* 343: 89-93 (1994)).

Although some details in the above two studies were recently questioned there is over-all agreement that PTP1B and SHP-1 play significant functional roles in platelet aggregation (Ezumi *et al.*, *J. Biol. Chem.* 270: 11927-11934 (1995)). In accordance with these observations, treatment of platelets with the PTPase inhibitor pervanadate leads to significant increase in tyrosine phosphorylation, secretion and aggregation (Pumiglia *et al.*, *Biochem. J.* 286: 441-449 (1992)).

15 PTPases: osteoporosis

The rate of bone formation is determined by the number and the activity of osteoblasts, which in turn are determined by the rate of proliferation and differentiation of osteoblast progenitor cells, respectively. Histomorphometric studies indicate that the osteoblast number is the primary determinant of the rate of bone formation in humans (Gruber *et al.*, *Mineral Electrolyte Metab.* 12: 246-254 (1987); reviewed in Lau *et al.*, *Biochem. J.* 257: 23-36 (1989)). Acid phosphatases/PTPases may be involved in negative regulation of osteoblast proliferation. Thus, fluoride, which has phosphatase inhibitory activity, has been found to increase spinal bone density in osteoporotics by increasing osteoblast proliferation (Lau *et al.*, *supra*). Consistent with this observation, an osteoblastic acid phosphatase with PTPase activity was found to be highly sensitive to mitogenic concentrations of fluoride (Lau *et al.*, *J. Biol. Chem.* 260: 4653-4660 (1985); Lau *et al.*, *J. Biol. Chem.* 262: 1389-1397 (1987); Lau *et al.*, *Adv. Protein Phosphatases* 4: 165-198 (1987)). Interestingly, it was recently found that the level of membrane-bound PTPase activity was increased dramatically when the

osteoblast-like cell line UMR 106.06 was grown on collagen type-I matrix compared to uncoated tissue culture plates. Since a significant increase in PTPase activity was observed in density-dependent growth arrested fibroblasts (Pallen and Tong, *Proc. Natl. Acad. Sci.* 88: 6996-7000 (1991)), it might be speculated that the increased PTPase activity directly inhibits cell growth. The mitogenic action of fluoride and other phosphatase inhibitors (molybdate and vanadate) may thus be explained by their inhibition of acid phosphatases/PTPases that negatively regulate the cell proliferation of osteoblasts. The complex nature of the involvement of PTPases in bone formation is further suggested by the recent identification of a novel parathyroid regulated, receptor-like PTPase, OST-PTP, expressed in bone and testis (Mauro *et al.*, *J. Biol. Chem.* 269: 30659-30667 (1994)). OST-PTP is up-regulated following differentiation and matrix formation of primary osteoblasts and subsequently down-regulated in the osteoblasts which are actively mineralizing bone in culture. It may be hypothesized that PTPase inhibitors may prevent differentiation via inhibition of OST-PTP or other PTPases thereby leading to continued proliferation. This would be in agreement with the above-mentioned effects of fluoride and the observation that the tyrosine phosphatase inhibitor orthovanadate appears to enhance osteoblast proliferation and matrix formation (Lau *et al.*, *Endocrinology* 116: 2463-2468 (1988)). In addition, it was recently observed that vanadate, vanadyl and pervanadate all increased the growth of the osteoblast-like cell line UMR106. Vanadyl and pervanadate were stronger stimulators of cell growth than vanadate. Only vanadate was able to regulate the cell differentiation as measured by cell alkaline phosphatase activity (Cortizo *et al.*, *Mol. Cell. Biochem.* 145: 97-102 (1995)).

PTPases: microorganisms

Dixon and coworkers have called attention to the fact that PTPases may be a key element in the pathogenic properties of *Yersinia* (reviewed in Clemens *et al. Molecular Microbiology* 5: 2617-2620 (1991)). This finding was rather surprising

since tyrosine phosphate is thought to be absent in bacteria. The genus *Yersinia* comprises 3 species: *Y. pestis* (responsible for the bubonic plague), *Y. pseudotuberculosis* and *Y. enterocolitica* (causing enteritis and mesenteric lymphadenitis). Interestingly, a dual-specificity phosphatase, VH1, has been identified in *Vaccinia virus* (Guan *et al.*, *Nature* **350**: 359-263 (1991)). These observations indicate that PTPases may play critical roles in microbial and parasitic infections, and they further point to PTPase inhibitors as a novel, putative treatment principle of infectious diseases.

10 SUMMARY OF THE INVENTION

The present invention relates to compounds of the general formula I, wherein A, R₁, R₂, R₃, R₄, R₁₆ and R₁₇ are as defined in the detailed part of the present description, wherein such compounds are pharmacologically useful inhibitors of Protein Tyrosine Phosphatases (PTPases) such as PTP1B, CD45, SHP-1, SHP-2, PTP α , LAR and HePTP or the like.

The present compounds are useful for the treatment, prevention, elimination, alleviation or amelioration of an indication related to type I diabetes, type II diabetes, impaired glucose tolerance, insulin resistance, obesity, immune dysfunctions including autoimmunity and AIDS, diseases with dysfunctions of the coagulation system, allergic diseases including asthma, osteoporosis, proliferative disorders including cancer and psoriasis, diseases with decreased or increased synthesis or effects of growth hormone, diseases with decreased or increased synthesis of hormones or cytokines that regulate the release of/or response to growth hormone, diseases of the brain including Alzheimer's disease and schizophrenia, and infectious diseases.

In another aspect, the present invention includes within its scope pharmaceutical compositions comprising, as an active ingredient, at least one of the compounds of the general formula I or a pharmaceutically acceptable salt thereof together with a pharmaceutically acceptable carrier or diluent.

In another aspect of the present invention there is provided a method of treating type I diabetes, type II diabetes, impaired glucose tolerance, insulin resistance, obesity, immune dysfunctions including autoimmunity and AIDS, diseases with dysfunctions of the coagulation system, allergic diseases including asthma, osteoporosis, proliferative disorders including cancer and psoriasis, diseases with decreased or increased synthesis or effects of growth hormone, diseases with decreased or increased synthesis of hormones or cytokines that regulate the release of/or response to growth hormone, diseases of the brain including Alzheimer's disease and schizophrenia, and infectious diseases.

10

The method of treatment may be described as the treatment, prevention, elimination, alleviation or amelioration of one of the above indications, which comprises the step of administering to the said subject a neurologically effective amount of a compound of the invention, or a pharmaceutically acceptable salt thereof.

15

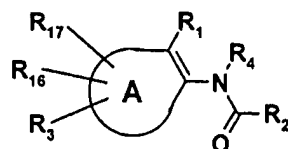
A further aspect of the invention relates to the use of a compound of the present invention for the preparation of a pharmaceutical composition for the treatment of all type I diabetes, type II diabetes, impaired glucose tolerance, insulin resistance, obesity, immune dysfunctions including autoimmunity and AIDS, diseases with dysfunctions of the coagulation system, allergic diseases including asthma, osteoporosis, proliferative disorders including cancer and psoriasis, diseases with decreased or increased synthesis or effects of growth hormone, diseases with decreased or increased synthesis of hormones or cytokines that regulate the release of/or response to growth hormone, diseases of the brain including Alzheimer's disease and schizophrenia, and infectious diseases.

20
25

DESCRIPTION OF THE INVENTION

The present invention relates to Compounds of the Formula 1 wherein A, R₁, R₂, R₃, R₄, R₁₆ and R₁₇ are defined below;

30



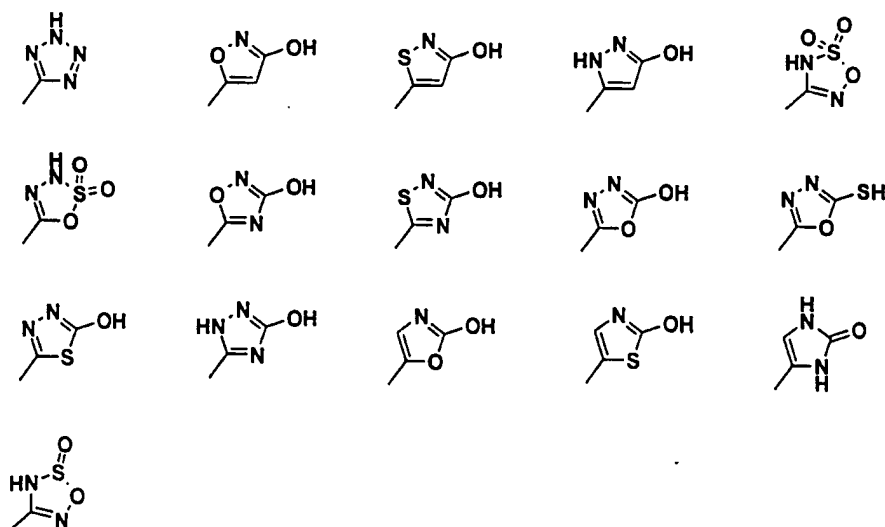
Formula 1

5

In the above Formula 1

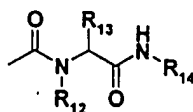
A is together with the double bond in Formula 1 phenyl, biphenyl, indenyl, fluorenyl, fluorenyl-9-one, naphthyl, pyridyl, pyridazinyl, pyrimidyl or pyrazinyl;

- 10 R_1 is hydrogen, COR_5 , OR_6 , CF_3 , nitro, cyano, SO_3H , $SO_2NR_7R_8$, $PO(OH)_2$, $CH_2PO(OH)_2$, $CHFPO(OH)_2$, $CF_2PO(OH)_2$, $C(=NH)NH_2$, NR_7R_8 or from the following 5-membered heterocycles:



15

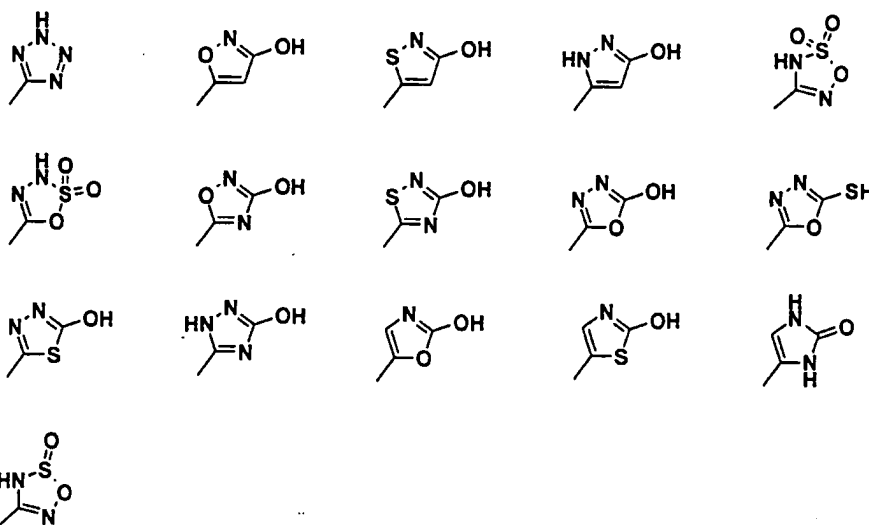
or R_1 is



wherein R_{12} , R_{13} , and R_{14} are independently hydrogen, C_1 - C_6 alkyl, aryl, aryl C_1 - C_6 alkyl and the alkyl and aryl groups are optionally substituted;

5

R_2 is COR_5 , OR_6 , CF_3 , nitro, cyano, SO_3H , $SO_2NR_7R_8$, $PO(OH)_2$, $CH_2PO(OH)_2$, $CHFPO(OH)_2$, $CF_2PO(OH)_2$, $C(=NH)NH_2$, NR_7R_8 , or selected from the following 5-membered heterocycles:

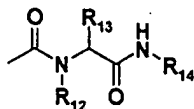


10

R_3 , R_{16} and R_{17} are independently hydrogen, halo, nitro, cyano, trihalomethyl, C_1 - C_6 alkyl, aryl, aryl C_1 - C_6 alkyl, hydroxy, carboxy, carboxy C_1 - C_6 alkyl, C_1 - C_6 alkyloxycarbonyl, aryloxycarbonyl, aryl C_1 - C_6 alkyloxycarbonyl, C_1 - C_6 alkyloxy, C_1 - C_6 alkyloxy C_1 - C_6 alkyl, aryloxy, aryl C_1 - C_6 alkyloxy, aryl C_1 - C_6 alkyloxy C_1 - C_6 alkyl, thio, C_1 - C_6 alkylthio, C_1 - C_6 alkylthio C_1 - C_6 alkyl, arylthio, aryl C_1 - C_6 alkylthio, aryl C_1 - C_6 alkylthio C_1 - C_6 alkyl, NR_7R_8 , C_1 - C_6 alkylamino C_1 - C_6 alkyl, aryl C_1 - C_6 alkylamino C_1 - C_6 alkyl, di(aryl C_1 - C_6 alkyl)amino C_1 - C_6 alkyl, C_1 - C_6 alkylcarbonyl, C_1 - C_6 alkylcarbonyl- C_1 - C_6 alkyl, aryl C_1 - C_6 alkylcarbonyl, aryl C_1 - C_6 alkylcarbonyl C_1 - C_6 alkyl, C_1 - C_6 alkylcarboxy, C_1 - C_6 alkylcarboxy C_1 - C_6 alkyl, arylcarboxy, arylcarboxy C_1 - C_6 alkyl, aryl C_1 - C_6 alkylcarboxy, aryl C_1 - C_6 alkylcarboxy C_1 - C_6 alkyl, C_1 - C_6 alkyl-carbonylamino, C_1 - C_6 alkylcarbonylamino C_1 - C_6 alkyl, -carbonyl NR_7C_1 - C_6 alkyl COR_{11} , aryl C_1 -

C₆alkylcarbonylamino, arylC₁-C₆alkylcarbonyl-aminoC₁-C₆alkyl, CONR₇R₈, or C₁-C₆alkylCONR₇R₈ wherein the alkyl and aryl groups are optionally substituted and R₁₁ is NR₇R₈, or C₁-C₆alkyl-NR₇R₈; or R₃ is

5



wherein R₁₂, R₁₃, and R₁₄ are independently hydrogen, C₁-C₆alkyl, aryl, arylC₁-C₆alkyl
10 and the alkyl and aryl groups are optionally substituted;

R₄ is hydrogen, hydroxy, C₁-C₆alkyl, aryl, arylC₁-C₆alkyl, NR₇R₈, C₁-C₆alkyloxy; whe-
rein the alkyl and aryl groups are optionally substituted;

15 R₅ is hydroxy, C₁-C₆alkyl, aryl, arylC₁-C₆alkyl, C₁-C₆alkyloxy, C₁-C₆alkyl-oxyC₁-
C₆alkyloxy, aryloxy, arylC₁-C₆alkyloxy, CF₃, NR₇R₈; wherein the alkyl and aryl groups
are optionally substituted;

R₆ is hydrogen, C₁-C₆alkyl, aryl, arylC₁-C₆alkyl; wherein the alkyl and aryl groups are
20 optionally substituted;

R₇ and R₈ are independently selected from hydrogen, C₁-C₆alkyl, aryl, arylC₁-C₆alkyl,
C₁-C₆alkylcarbonyl, arylcarbonyl, arylC₁-C₆alkylcarbonyl, C₁-C₆alkylcarboxy or arylC₁-
C₆alkylcarboxy wherein the alkyl and aryl groups are optionally substituted; or
25 R₇ and R₈ are together with the nitrogen to which they are attached forming a satu-
rated, partially saturated or aromatic cyclic, bicyclic or tricyclic ring system containing
from 3 to 14 carbon atoms and from 0 to 3 additional heteroatoms selected from ni-
trogen, oxygen or sulfur, the ring system can optionally be substituted with at least
one C₁-C₆alkyl, aryl, arylC₁-C₆alkyl, hydroxy, oxo, C₁-C₆alkyloxy, arylC₁-C₆alkyloxy,
30 C₁-C₆alkyloxyC₁-C₆alkyl, NR₉R₁₀ or C₁-C₆alkylaminoC₁-C₆alkyl, wherein R₉ and R₁₀ are
independently selected from hydrogen, C₁-C₆alkyl, aryl, arylC₁-C₆alkyl, C₁-

C₆alkylcarbonyl, arylcarbonyl, arylC₁-C₆alkylcarbonyl, C₁-C₆alkylcarboxy or arylC₁-C₆alkylcarboxy; wherein the alkyl and aryl groups are optionally substituted; or R₇ and R₈ are independently a saturated or partial saturated cyclic 5, 6 or 7 membered amine, imide or lactam;

5

or a salt thereof with a pharmaceutically acceptable acid or base, or any optical isomer or mixture of optical isomers, including a racemic mixture, or any tautomeric forms.

10 DEFINITIONS

Signal transduction is a collective term used to define all cellular processes that follow the activation of a given cell or tissue. Examples of signal transduction, which are not intended to be in any way limiting to the scope of the invention claimed, are cellular events that are induced by polypeptide hormones and growth factors (e.g. insulin, insulin-like growth factors I and II, growth hormone, epidermal growth factor, platelet-derived growth factor), cytokines (e.g. interleukins), extracellular matrix components, and cell-cell interactions.

Phosphotyrosine recognition units/tyrosine phosphate recognition units/pTyr recognition units are defined as areas or domains of proteins or glycoproteins that have affinity for molecules containing phosphorylated tyrosine residues (pTyr). Examples of pTyr recognition units, which are not intended to be in any way limiting to the scope of the invention claimed, are: PTPases, SH2 domains and PTB domains.

PTPases are defined as enzymes with the capacity to dephosphorylate pTyr-containing proteins or glycoproteins. Examples of PTPases, which are not intended to be in any way limiting to the scope of the invention claimed, are: 'classical' PTPases (intracellular PTPases (e.g. PTP1B, TC-PTP, PTP1C, PTP1D, PTPD1, PTPD2) and

30

receptor-type PTPases (e.g. PTP α , PTP ϵ , PTP β , PTP γ , CD45, PTP κ , PTP μ), dual specificity phosphatases (VH1, VHR, cdc25), LMW-PTPases or acid phosphatases.

SH2 domains (Src homology 2 domains) are non-catalytic protein modules that bind to pTyr (phosphotyrosine residue) containing proteins, i.e. SH2 domains are pTyr recognition units. SH2 domains, which consist of ~100 amino acid residues, are found in a number of different molecules involved in signal transduction processes. The following is a non-limiting list of proteins containing SH2 domains: Src, Hck, Lck, Syk, Zap70, SHP-1, SHP-2, STATs, Grb-2, Shc, p85/PI3K, Gap, vav (see Russell et al, FEBS Lett. 304:15-20 (1992); Pawson, Nature 373: 573-580 (1995); Sawyer, Biopolymers (Peptide Science) 47: 243-261 (1998); and references herein).

As used herein, the term "attached" or "-" (e.g. -COR₁₁, which indicates the carbonyl attachment point to the scaffold) signifies a stable covalent bond, certain preferred points of attachment points being apparent to those skilled in the art. The terms "halogen" or "halo" include fluorine, chlorine, bromine, and iodine. The term "alkyl" includes C₁-C₆ straight chain saturated and C₂-C₆ unsaturated aliphatic hydrocarbon groups, C₁-C₆ branched saturated and C₂-C₆ unsaturated aliphatic hydrocarbon groups, C₃-C₆ cyclic saturated and C₅-C₆ unsaturated aliphatic hydrocarbon groups, and C₁-C₆ straight chain or branched saturated and C₂-C₆ straight chain or branched unsaturated aliphatic hydrocarbon groups substituted with C₃-C₆ cyclic saturated and unsaturated aliphatic hydrocarbon groups having the specified number of carbon atoms. For example, this definition shall include but is not limited to methyl (Me), ethyl (Et), propyl (Pr), butyl (Bu), pentyl, hexyl, heptyl, ethenyl, propenyl, butenyl, penentyl, hexenyl, isopropyl (i-Pr), isobutyl (i-Bu), *tert*-butyl (*t*-Bu), *sec*-butyl (*s*-Bu), isopentyl, neopentyl, cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cyclopentenyl, cyclohexenyl, methylcyclopropyl, ethylcyclohexenyl, butenylcyclopentyl, and the like.

The term "substituted alkyl" represents an alkyl group as defined above wherein the substituents are independently selected from halo, cyano, nitro, trihalomethyl,

carbamoyl, hydroxy, oxo, COR₅, C₁-C₆alkyl, C₁-C₆alkyloxy, aryloxy, arylC₁-C₆alkyloxy, thio, C₁-C₆alkylthio, arylthio, arylC₁-C₆alkylthio, NR₇R₈, C₁-C₆alkylamino, arylamino, arylC₁-C₆alkylamino, di(arylC₁-C₆alkyl)amino, C₁-C₆alkylcarbonyl, arylC₁-C₆alkylcarbonyl, C₁-C₆alkylcarboxy, arylcarboxy, arylC₁-C₆alkylcarboxy, C₁-C₆alkylcarbonyl-amino, -C₁-C₆alkylaminoCOR₁₁, arylC₁-C₆alkylcarbonylamino, tetrahydrofuranyl, morpholinyl, piperazinyl, -CONR₇R₈, -C₁-C₆alkyl-CONR₇R₈, or a saturated or partial saturated cyclic 5, 6 or 7 membered amine, imide or lactam; wherein R₁₁ is hydroxy, C₁-C₆alkyl, aryl, arylC₁-C₆alkyl, C₁-C₆alkyloxy, aryloxy, arylC₁-C₆alkyloxy and R₅ is defined as above or NR₇R₈, wherein R₇, R₈ are defined as above.

The term "saturated, partially saturated or aromatic cyclic, bicyclic or tricyclic ring system" represents but are not limit to aziridinyl, pyrrolyl, pyrrolinyl, pyrrolidinyl, imidazolyl, 2-imidazolinyl, imidazolidinyl, pyrazolyl, 2-pyrazolinyl, 1,2,3-triazolyl, 1,2,4-triazolyl, morpholinyl, piperidinyl, thiomorpholinyl, piperazinyl, indolyl, isoindolyl, 1,2,3,4-tetrahydro-quinolinyl, 1,2,3,4-tetrahydro-isoquinolinyl, 1,2,3,4-tetrahydro-quinoxalyl, indolyl, indazolyl, benzimidazolyl, benzotriazolyl, purinyl, carbazolyl, acridinyl, phenothiazinyl, phenoxazinyl, iminodibenzyl, iminostilbenyl.

The term "alkyloxy" (e.g. methoxy, ethoxy, propyloxy, allyloxy, cyclohexyloxy) represents an "alkyl" group as defined above having the indicated number of carbon atoms attached through an oxygen bridge. The term "alkyloxyalkyl" represents an "alkyloxy" group attached through an alkyl group as defined above having the indicated number of carbon atoms.

The term "alkyloxyalkyloxy" represents an "alkyloxyalkyl" group attached through an oxygen atom as defined above having the indicated number of carbon atoms.

The term "aryloxy" (e.g. phenoxy, naphthyloxy and the like) represents an aryl group as defined below attached through an oxygen bridge.

The term "arylalkyloxy" (e.g. phenethyloxy, naphthylmethyloxy and the like) represents an "arylalkyl" group as defined below attached through an oxygen bridge.

The term "arylalkyloxyalkyl" represents an "arylalkyloxy" group as defined above attached through an "alkyl" group defined above having the indicated number of carbon atoms.

5 The term "arylthio" (e.g. phenylthio, naphthylthio and the like) represents an "aryl" group as defined below attached through an sulfur bridge.

The term "alkyloxycarbonyl" (e.g. methylformiat, ethylformiat and the like) represents an "alkyloxy" group as defined above attached through a carbonyl group.

10 The term "aryloxycarbonyl" (e.g. phenylformiat, 2-thiazolylformiat and the like) represents an "aryloxy" group as defined above attached through a carbonyl group.

The term "arylalkyloxycarbonyl" (e.g. benzylformiat, phenylethylformiat and the like) represents an "arylalkyloxy" group as defined above attached through a carbonyl group.

15 The term "alkyloxycarbonylalkyl" represents an "alkyloxycarbonyl" group as defined above attached through an "alkyl" group as defined above having the indicated number of carbon atoms.

The term "arylalkyloxycarbonylalkyl" represents an "arylalkyloxycarbonyl" group as defined above attached through an "alkyl" group as defined above having the indicated number of carbon atoms.

20

The term "alkylthio" (e.g. methylthio, ethylthio, propylthio, cyclohexenylthio and the like) represents an "alkyl" group as defined above having the indicated number of carbon atoms attached through a sulfur bridge.

25 The term "arylalkylthio" (e.g. phenylmethylthio, phenylethylthio, and the like) represents an "arylalkyl" group as defined above having the indicated number of carbon atoms attached through a sulfur bridge.

The term "alkylthioalkyl" represents an "alkylthio" group attached through an alkyl group as defined above having the indicated number of carbon atoms.

30 The term "arylalkylthioalkyl" represents an "arylalkylthio" group attached through an alkyl group as defined above having the indicated number of carbon atoms.

The term "alkylamino" (e.g. methylamino, diethylamino, butylamino, N-propyl-N-hexylamino, (2-cyclopentyl)propylamino, hexenylamino, pyrrolidinyl, piperidinyl and the like) represents one or two "alkyl" groups as defined above having the indicated number of carbon atoms attached through an amine bridge. The two alkyl groups
5 may be taken together with the nitrogen to which they are attached forming a saturated, partially saturated or aromatic cyclic, bicyclic or tricyclic ring system containing from 3 to 14 carbon atoms and from 0 to 3 additional heteroatoms selected from nitrogen, oxygen or sulfur, the ring system can optionally be substituted with at least one C₁-C₆alkyl, aryl, arylC₁-C₆alkyl, hydroxy, oxo, C₁-C₆alkyloxy, arylC₁-C₆alkyloxy,
10 C₁-C₆alkyloxyC₁-C₆alkyl, NR₉R₁₀ or C₁-C₆alkylaminoC₁-C₆alkyl substituent wherein the alkyl and aryl groups are optionally substituted as defined in the definition section and R₉ and R₁₀ are defined as above.

The term "arylalkylamino" (e.g. benzylamino, diphenylethylamino and the like) represents one or two "arylalkyl" groups as defined above having the indicated number of
15 carbon atoms attached through an amine bridge. The two "arylalkyl" groups may be taken together with the nitrogen to which they are attached forming a saturated, partially saturated or aromatic cyclic, bicyclic or tricyclic ring system containing from 3 to 14 carbon atoms and from 0 to 3 additional heteroatoms selected from nitrogen, oxygen or sulfur, the ring system can optionally be substituted with at least one C₁-
20 C₆alkyl, aryl, arylC₁-C₆alkyl, hydroxy, oxo, C₁-C₆alkyloxy, arylC₁-C₆alkyloxy, C₁-C₆alkyloxyC₁-C₆alkyl, NR₉R₁₀ or C₁-C₆alkylaminoC₁-C₆alkyl substituent wherein the alkyl and aryl groups are optionally substituted as defined in the definition section and R₉ and R₁₀ are defined as above.

The term "alkylaminoalkyl" represents an "alkylamino" group attached through an alkyl group as defined above having the indicated number of carbon atoms.
25

The term "arylalkylaminoalkyl" represents an "arylalkylamino" group attached through an alkyl group as defined above having the indicated number of carbon atoms.

The term "arylalkyl" (e.g. benzyl, phenylethyl) represents an "aryl" group as defined below attached through an alkyl having the indicated number of carbon atoms or
30 substituted alkyl group as defined above.

The term "alkylcarbonyl" (e.g. cyclooctylcarbonyl, pentylcarbonyl, 3-hexenylcarbonyl) represents an "alkyl" group as defined above having the indicated number of carbon atoms attached through a carbonyl group.

5 The term "arylcarbonyl" (benzoyl) represents an "aryl" group as defined above attached through a carbonyl group.

The term "arylalkylcarbonyl" (e.g. phenylcyclopropylcarbonyl, phenylethylcarbonyl and the like) represents an "arylalkyl" group as defined above having the indicated number of carbon atoms attached through a carbonyl group.

10 The term "alkylcarbonylalkyl" represents an "alkylcarbonyl" group attached through an "alkyl" group as defined above having the indicated number of carbon atoms.

The term "arylalkylcarbonylalkyl" represents an "arylalkylcarbonyl" group attached through an alkyl group as defined above having the indicated number of carbon atoms.

15 The term "alkylcarboxy" (e.g. heptylcarboxy, cyclopropylcarboxy, 3-pentenylcarboxy) represents an "alkylcarbonyl" group as defined above wherein the carbonyl is in turn attached through an oxygen bridge.

20 The term "arylalkylcarboxy" (e.g. benzylcarboxy, phenylcyclopropylcarboxy and the like) represents an "arylalkylcarbonyl" group as defined above wherein the carbonyl is in turn attached through an oxygen bridge.

The term "arylcarboxyalkyl" (e.g. phenylcarboxymethyl) represents an "arylcarbonyl" group defined above wherein the carbonyl is in turn attached through an oxygen bridge to an alkyl chain having the indicated number of carbon atoms.

25 The term "alkylcarboxyalkyl" represents an "alkylcarboxy" group attached through an "alkyl" group as defined above having the indicated number of carbon atoms.

The term "arylalkylcarboxyalkyl" represents an "arylalkylcarboxy" group attached through an "alkyl" group as defined above having the indicated number of carbon atoms.

30 The term "alkylcarbonylamino" (e.g. hexylcarbonylamino, cyclopentylcarbonylaminomethyl, methylcarbonylaminophenyl) represents an "alkylcarbonyl" group as defined above wherein the carbonyl is in turn attached through the nitrogen atom of

an amino group. The nitrogen atom may itself be substituted with an alkyl or aryl group.

The term "arylalkylcarbonylamino" (e.g. benzylcarbonylamino and the like) represents an "arylalkylcarbonyl" group as defined above wherein the carbonyl is in turn
 5 attached through the nitrogen atom of an amino group. The nitrogen atom may itself be substituted with an alkyl or aryl group.

The term "alkylcarbonylaminoalkyl" represents an "alkylcarbonylamino" group attached through an "alkyl" group as defined above having the indicated number of carbon atoms. The nitrogen atom may itself be substituted with an alkyl or aryl group.

10 The term "arylalkylcarbonylaminoalkyl" represents an "arylalkylcarbonylamino" group attached through an "alkyl" group as defined above having the indicated number of carbon atoms. The nitrogen atom may itself be substituted with an alkyl or aryl group.

The term "alkylcarbonylaminoalkylcarbonyl" represents an alkylcarbonylaminoalkyl
 15 group attached through a carbonyl group. The nitrogen atom may be further substituted with an "alkyl" or "aryl" group.

The term "aryl" represents an unsubstituted, mono-, di- or trisubstituted monocyclic, polycyclic, biaryl and heterocyclic aromatic groups covalently attached at any ring
 20 position capable of forming a stable covalent bond, certain preferred points of attachment being apparent to those skilled in the art (e.g., 3-indolyl, 4-imidazolyl). The aryl substituents are independently selected from the group consisting of halo, nitro, cyano, trihalomethyl, C₁-C₆alkyl, aryl, arylC₁-C₆alkyl, hydroxy, COR₅, C₁-C₆alkyloxy, C₁-C₆alkyloxyC₁-C₆alkyl, aryloxy, arylC₁-C₆alkyloxy, arylC₁-C₆alkyloxyC₁-C₆alkyl, thio,
 25 C₁-C₆alkylthio, C₁-C₆alkylthioC₁-C₆alkyl, arylthio, arylC₁-C₆alkylthio, arylC₁-C₆alkylthioC₁-C₆alkyl, NR₂R₃, C₁-C₆alkylamino, C₁-C₆alkylaminoC₁-C₆alkyl, arylamino, arylC₁-C₆alkylamino, arylC₁-C₆alkyl-aminoC₁-C₆alkyl, di(arylC₁-C₆alkyl)aminoC₁-C₆alkyl, C₁-C₆alkylcarbonyl, C₁-C₆alkylcarbonylC₁-C₆alkyl, arylC₁-C₆alkylcarbonyl, arylC₁-C₆alkyl-carbonylC₁-C₆alkyl, C₁-C₆alkylcarboxy, C₁-C₆alkylcarboxyC₁-C₆alkyl, arylC₁-C₆alkylcarboxy, arylC₁-C₆alkylcarboxyC₁-C₆alkyl, carboxyC₁-C₆alkyloxy, C₁-C₆alkylcarbonylamino, C₁-C₆alkylcarbonyl-aminoC₁-C₆alkyl, -carbonylNR₂C₁-
 30

C₆alkylCOR₁₁, arylC₁-C₆alkylcarbonyl-amino, arylC₁-C₆alkylcarbonylaminoC₁-C₆alkyl, -CONR₈R₉, or -C₁-C₆alkyl-CONR₈R₉; wherein R₇, R₈, R₉, and R₁₁ are defined as above and the alkyl and aryl groups are optionally substituted as defined in the definition section;

5

The definition of aryl includes but is not limited to phenyl, biphenyl, indenyl, fluorenyl, naphthyl (1-naphthyl, 2-naphthyl), pyrrolyl (2-pyrrolyl), pyrazolyl (3-pyrazolyl), imidazolyl (1-imidazolyl, 2-imidazolyl, 4-imidazolyl, 5-imidazolyl), triazolyl (1,2,3-triazol-1-yl, 1,2,3-triazol-2-yl, 1,2,3-triazol-4-yl, 1,2,4-triazol-3-yl), oxazolyl (2-oxazolyl, 4-oxazolyl, 5-oxazolyl), isoxazolyl (3-isoxazolyl, 4-isoxazolyl, 5-isoxazolyl), thiazolyl (2-thiazolyl, 4-thiazolyl, 5-thiazolyl), thiophenyl (2-thiophenyl, 3-thiophenyl, 4-thiophenyl, 5-thiophenyl), furanyl (2-furanyl, 3-furanyl, 4-furanyl, 5-furanyl), pyridyl (2-pyridyl, 3-pyridyl, 4-pyridyl, 5-pyridyl), 5-tetrazolyl, pyrimidinyl (2-pyrimidinyl, 4-pyrimidinyl, 5-pyrimidinyl, 6-pyrimidinyl), pyrazinyl, pyridazinyl (3-pyridazinyl, 4-pyridazinyl, 5-pyridazinyl), quinolyl (2-quinolyl, 3-quinolyl, 4-quinolyl, 5-quinolyl, 6-quinolyl, 7-quinolyl, 8-quinolyl), isoquinolyl (1-isoquinolyl, 3-isoquinolyl, 4-isoquinolyl, 5-isoquinolyl, 6-isoquinolyl, 7-isoquinolyl, 8-isoquinolyl), benzo[b]furanyl (2-benzo[b]furanyl, 3-benzo[b]furanyl, 4-benzo[b]furanyl, 5-benzo[b]furanyl, 6-benzo[b]furanyl, 7-benzo[b]furanyl), 2,3-dihydro-benzo[b]furanyl (2-(2,3-dihydro-benzo[b]furanyl), 3-(2,3-dihydro-benzo[b]furanyl), 4-(2,3-dihydro-benzo[b]furanyl), 5-(2,3-dihydro-benzo[b]furanyl), 6-(2,3-dihydro-benzo[b]furanyl), 7-(2,3-dihydro-benzo[b]furanyl)), benzo[b]thiophenyl (2-benzo[b]thiophenyl, 3-benzo[b]thiophenyl, 4-benzo[b]thiophenyl, 5-benzo[b]thiophenyl, 6-benzo[b]thiophenyl, 7-benzo[b]thiophenyl), 2,3-dihydro-benzo[b]-thiophenyl (2-(2,3-dihydro-benzo[b]thiophenyl), 3-(2,3-dihydro-benzo[b]-thiophenyl), 4-(2,3-dihydro-benzo[b]thiophenyl), 5-(2,3-dihydro-benzo[b]-thiophenyl), 6-(2,3-dihydro-benzo[b]thiophenyl), 7-(2,3-dihydro-benzo[b]-thiophenyl)), 4,5,6,7-tetrahydro-benzo[b]thiophenyl (2-(4,5,6,7-tetrahydro-benzo[b]thiophenyl), 3-(4,5,6,7-tetrahydro-benzo[b]thiophenyl), 4-(4,5,6,7-tetrahydro-benzo[b]thiophenyl), 5-(4,5,6,7-tetrahydro-benzo[b]thiophenyl), 6-(4,5,6,7-tetrahydro-benzo[b]thiophenyl), 7-(4,5,6,7-tetrahydro-benzo[b]thiophenyl)), 4,5,6,7-tetrahydro-thieno[2,3-c]pyridyl (4-

(4,5,6,7-tetrahydro-thieno[2,3-c]pyridyl), 5-(4,5,6,7-tetrahydro-thieno[2,3-c]pyridyl), 6-(4,5,6,7-tetrahydro-thieno[2,3-c]pyridyl), 7-(4,5,6,7-tetrahydro-thieno[2,3-c]pyridyl)), indolyl (1-indolyl, 2-indolyl, 3-indolyl, 4-indolyl, 5-indolyl, 6-indolyl, 7-indolyl), isoindolyl (1-isoindolyl, 2-isoindolyl, 3-isoindolyl, 4-isoindolyl, 5-isoindolyl, 6-isoindolyl, 7-isoindolyl), 1,3-dihydro-isoindolyl (1-(1,3-dihydro-isoindolyl), 2-(1,3-dihydro-isoindolyl), 3-(1,3-dihydro-isoindolyl), 4-(1,3-dihydro-isoindolyl), 5-(1,3-dihydro-isoindolyl), 6-(1,3-dihydro-isoindolyl), 7-(1,3-dihydro-isoindolyl)), indazole (1-indazolyl, 3-indazolyl, 4-indazolyl, 5-indazolyl, 6-indazolyl, 7-indazolyl), benzimidazolyl (1-benzimidazolyl, 2-benzimidazolyl, 4-benzimidazolyl, 5-benzimidazolyl, 6-benzimidazolyl, 7-benzimidazolyl, 8-benzimidazolyl), benzoxazolyl (1-benz-oxazolyl, 2-benzoxazolyl), benzothiazolyl (1-benzothiazolyl, 2-benzothiazolyl, 4-benzothiazolyl, 5-benzothiazolyl, 6-benzothiazolyl, 7-benzothiazolyl), carbazolyl (1-carbazolyl, 2-carbazolyl, 3-carbazolyl, 4-carbazolyl), 5H-dibenz[b,f]azepine (5H-dibenz[b,f]azepin-1-yl, 5H-dibenz[b,f]azepine-2-yl, 5H-dibenz[b,f]azepine-3-yl, 5H-dibenz[b,f]azepine-4-yl, 5H-dibenz[b,f]azepine-5-yl), 10,11-dihydro-5H-dibenz[b,f]azepine (10,11-dihydro-5H-dibenz[b,f]azepine-1-yl, 10,11-dihydro-5H-dibenz[b,f]azepine-2-yl, 10,11-dihydro-5H-dibenz[b,f]azepine-3-yl, 10,11-dihydro-5H-dibenz[b,f]azepine-4-yl, 10,11-dihydro-5H-dibenz[b,f]azepine-5-yl), piperidinyl (2-piperidinyl, 3-piperidinyl, 4-piperidinyl), pyrrolidinyl (1-pyrrolidinyl, 2-pyrrolidinyl, 3-pyrrolidinyl), phenylpyridyl (2-phenyl-pyridyl, 3-phenyl-pyridyl, 4-phenylpyridyl), phenylpyrimidinyl (2-phenylpyrimidinyl, 4-phenyl-pyrimidinyl, 5-phenylpyrimidinyl, 6-phenylpyrimidinyl), phenylpyrazinyl, phenylpyridazinyl (3-phenylpyridazinyl, 4-phenylpyridazinyl, 5-phenyl-pyridazinyl).

25 The term "arylcarbonyl" (e.g. 2-thiophenylcarbonyl, 3-methoxy-anthrylcarbonyl, oxazolylcarbonyl) represents an "aryl" group as defined above attached through a carbonyl group.

The term "arylalkylcarbonyl" (e.g. (2,3-dimethoxyphenyl)-propylcarbonyl, (2-chloronaphthyl)pentenylcarbonyl, imidazolylcyclopentylcarbonyl) represents an
30 "arylalkyl" group as defined above wherein the "alkyl" group is in turn attached through a carbonyl.

The compounds of the present invention have asymmetric centers and may occur as racemates, racemic mixtures, and as individual enantiomers or diastereoisomers, with all isomeric forms being included in the present invention as well as mixtures thereof.

5

Pharmaceutically acceptable salts of the compounds of formula 1, where a basic or acidic group is present in the structure, are also included within the scope of this invention. When an acidic substituent is present, such as -COOH , 5-tetrazolyl or -P(O)(OH)_2 , there can be formed the ammonium, morpholinium, sodium, potassium, barium, calcium salt, and the like, for use as the dosage form. When a basic group is present, such as amino or a basic heteroaryl radical, such as pyridyl, an acidic salt, such as hydrochloride, hydrobromide, phosphate, sulfate, trifluoroacetate, trichloroacetate, acetate, oxalate, maleate, pyruvate, malonate, succinate, citrate, tartarate, fumarate, mandelate, benzoate, cinnamate, methanesulfonate, ethane sulfonate, picroate and the like, and include acids related to the pharmaceutically acceptable salts listed in Journal of Pharmaceutical Science, 66, 2 (1977) and incorporated herein by reference, can be used as the dosage form.

Also, in the case of the -COOH or -P(O)(OH)_2 being present, pharmaceutically acceptable esters can be employed, e.g., methyl, tert-butyl, acetoxymethyl, pivaloyloxymethyl, and the like, and those esters known in the art for modifying solubility or hydrolysis characteristics for use as sustained release or prodrug formulations.

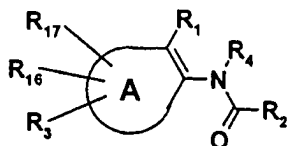
In addition, some of the compounds of the instant invention may form solvates with water or common organic solvents. Such solvates are encompassed within the scope of the invention.

The term "therapeutically effective amount" shall mean that amount of drug or pharmaceutical agent that will elicit the biological or medical response of a tissue, system, animal, or human that is being sought by a researcher, veterinarian, medical doctor or other.

30

PREFERRED EMBODIMENTS OF THE INVENTION

- 5 Compounds of Formula 1a are preferred compounds of the invention



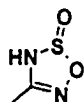
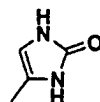
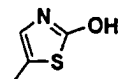
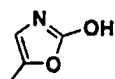
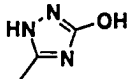
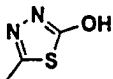
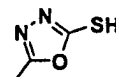
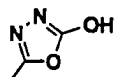
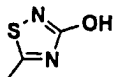
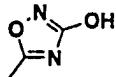
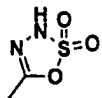
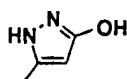
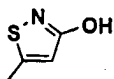
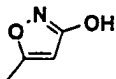
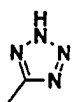
Formula 1a

10

wherein

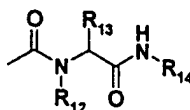
A is together with the double bond in Formula 1a phenyl, biphenyl, indenyl, fluorenyl, fluorenyl-9-one or naphthyl;

- 15 R_1 is COR_5 , OR_6 , CF_3 , nitro, cyano, SO_3H , $SO_2NR_7R_8$, $PO(OH)_2$, $CH_2PO(OH)_2$, $CHFPO(OH)_2$, $CF_2PO(OH)_2$, $C(=NH)NH_2$, NR_7R_8 or selected from the following 5-membered heterocycles:



or R_1 is

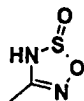
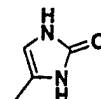
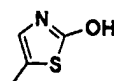
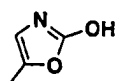
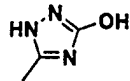
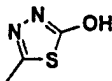
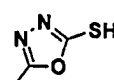
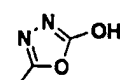
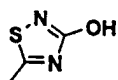
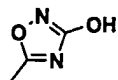
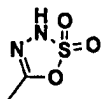
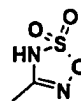
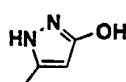
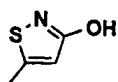
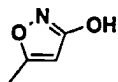
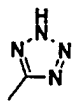
5



wherein R_{12} , R_{13} , and R_{14} are independently hydrogen, C_1 - C_6 alkyl, aryl, aryl C_1 - C_6 alkyl and the alkyl and aryl groups are optionally substituted;

10

R_2 is COR_5 , OR_6 , CF_3 , nitro, cyano, SO_3H , $SO_2NR_7R_8$, $PO(OH)_2$, $CH_2PO(OH)_2$, $CHFPO(OH)_2$, $CF_2PO(OH)_2$, $C(=NH)NH_2$, NR_7R_8 , or selected from the following 5-membered heterocycles:



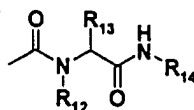
15

R_3 , R_{16} and R_{17} are independently hydrogen, halo, nitro, cyano, trihalomethyl, C_1 - C_6 alkyl, aryl, aryl C_1 - C_6 alkyl, hydroxy, carboxy, carboxy C_1 - C_6 alkyl, C_1 - C_6 alkyloxy-carbonyl, aryloxy-carbonyl, aryl C_1 - C_6 alkyloxy-carbonyl, C_1 - C_6 alkyloxy, C_1 - C_6 alkyloxy C_1 - C_6 alkyl, aryloxy, aryl C_1 - C_6 alkyloxy, aryl C_1 - C_6 alkyl-oxy C_1 - C_6 alkyl, thio, C_1 - C_6 alkylthio, C_1 - C_6 alkylthio C_1 - C_6 alkyl, arylthio, aryl C_1 - C_6 alkylthio, aryl C_1 - C_6 alkylthio C_1 - C_6 alkyl, NR_7R_8 , C_1 - C_6 alkyl-amino C_1 - C_6 alkyl, aryl C_1 - C_6 alkylamino C_1 - C_6 alkyl, di(aryl C_1 -

20

C_6 alkyl)-amino C_1 - C_6 alkyl, C_1 - C_6 alkylcarbonyl, C_1 - C_6 alkylcarbonyl C_1 - C_6 alkyl, aryl C_1 - C_6 alkylcarbonyl, aryl C_1 - C_6 alkylcarbonyl C_1 - C_6 alkyl, C_1 - C_6 alkyl-carboxy, C_1 - C_6 alkyl-carboxy C_1 - C_6 alkyl, arylcarboxy, aryl C_1 - C_6 alkyl-carboxy, aryl C_1 - C_6 alkylcarboxy C_1 - C_6 alkyl, C_1 - C_6 alkylcarbonylamino, C_1 - C_6 alkylcarbonyl-amino C_1 - C_6 alkyl,
 5 -carbonyl NR_7C_1 - C_6 alkylCOR₁₁, aryl C_1 - C_6 alkyl-carbonylamino, aryl C_1 - C_6 alkylcarbonylamino C_1 - C_6 alkyl, CONR₇R₈, or C_1 - C_6 alkylCONR₇R₈ wherein the alkyl and aryl groups are optionally substituted and R₁₁ is NR₇R₈, or C_1 - C_6 alkylNR₇R₈; or R₃ is

10



15

wherein R₁₂, R₁₃, and R₁₄ are independently hydrogen, C_1 - C_6 alkyl, aryl, aryl C_1 - C_6 alkyl and the alkyl and aryl groups are optionally substituted;

R₄ is hydrogen, hydroxy, C_1 - C_6 alkyl, aryl, aryl C_1 - C_6 alkyl, NR₇R₈, C_1 - C_6 alkyloxy; wherein the alkyl and aryl groups are optionally substituted;

20

R₅ is hydroxy, C_1 - C_6 alkyl, aryl, aryl C_1 - C_6 alkyl, CF₃, NR₇R₈; wherein the alkyl and aryl groups are optionally substituted;

R₆ is hydrogen, C_1 - C_6 alkyl, aryl, aryl C_1 - C_6 alkyl; wherein the alkyl and aryl groups are optionally substituted;

25

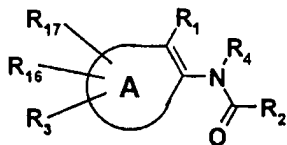
R₇ and R₈ are independently selected from hydrogen, C_1 - C_6 alkyl, aryl, aryl C_1 - C_6 alkyl, C_1 - C_6 alkyl-carbonyl, arylcarbonyl, aryl C_1 - C_6 alkyl-carbonyl, C_1 - C_6 alkyl-carboxy or aryl C_1 - C_6 alkylcarboxy wherein the alkyl and aryl groups are optionally substituted; or R₇ and R₈ are taken together with the nitrogen to which they are attached forming a
 30 cyclic or bicyclic system containing 3 to 11 carbon atoms and 0 to 2 additional heteroatoms selected from nitrogen, oxygen or sulfur, the ring system can optionally be substituted with at least one C_1 - C_6 alkyl, aryl, aryl C_1 - C_6 alkyl, hydroxy, C_1 - C_6 alkyloxy,

arylC₁-C₆alkoxy, C₁-C₆alkoxyC₁-C₆alkyl, NR₉R₁₀ or C₁-C₆alkylamino-C₁-C₆alkyl, wherein R₉ and R₁₀ are independently selected from hydrogen, C₁-C₆alkyl, aryl, arylC₁-C₆alkyl, C₁-C₆alkylcarbonyl, arylcarbonyl, arylC₁-C₆alkylcarbonyl, C₁-C₆alkylcarboxy or arylC₁-C₆alkylcarboxy; wherein the alkyl and aryl groups are optionally substituted; or

R₇ and R₈ are independently a saturated or partial saturated cyclic 5, 6 or 7 membered amine or lactam;

or a salt thereof with a pharmaceutically acceptable acid or base, or any optical isomer or mixture of optical isomers, including a racemic mixture, or any tautomeric forms.

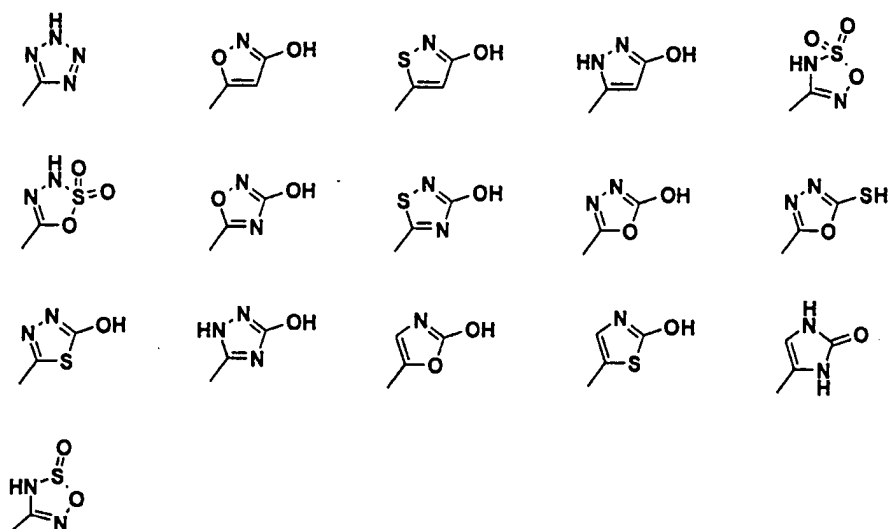
Also, compounds of Formula 1c are preferred compounds of the invention



Formula 1c

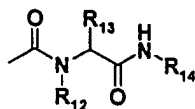
wherein A is together with the double bond in Formula 1c pyridyl, pyridazinyl, pyrimidyl or pyrazinyl;

R₁ is COR₅, OR₆, CF₃, nitro, cyano, SO₃H, SO₂NR₇R₈, PO(OH)₂, CH₂PO(OH)₂, CHFPO(OH)₂, CF₂PO(OH)₂, C(=NH)NH₂, NR₇R₈ or selected from the following 5-membered heterocycles:



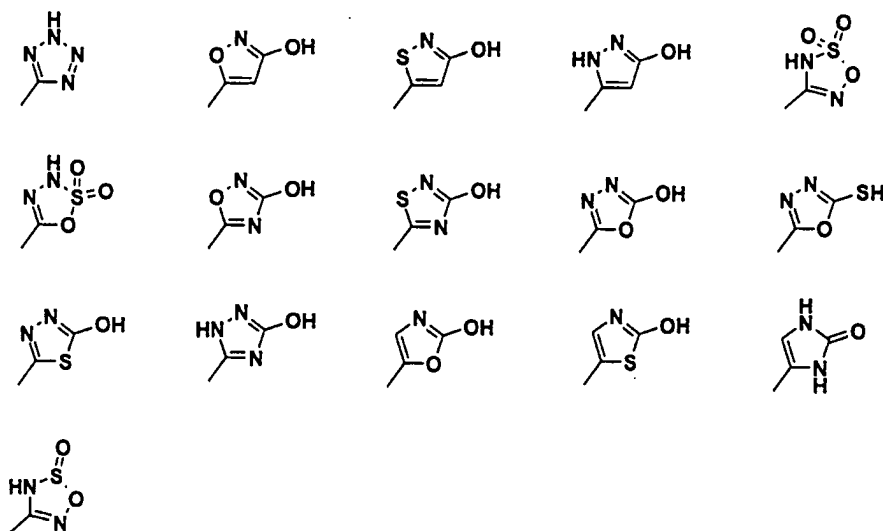
or R_1 is

5

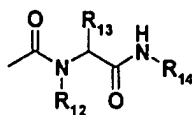


wherein R_{12} , R_{13} , and R_{14} are independently hydrogen, C_1 - C_6 alkyl, aryl,
 10 aryl C_1 - C_6 alkyl and the alkyl and aryl groups are optionally substituted;

R_2 is COR_5 , OR_6 , CF_3 , nitro, cyano, SO_3H , $SO_2NR_7R_8$, $PO(OH)_2$, $CH_2PO(OH)_2$,
 $CHFPO(OH)_2$, $CF_2PO(OH)_2$, $C(=NH)NH_2$, NR_7R_8 , or selected from the following 5-
 15 membered heterocycles:



- R_3 , R_{16} and R_{17} are independently hydrogen, halo, nitro, cyano, trihalomethyl, C_1 - C_6 alkyl, aryl, aryl C_1 - C_6 alkyl, hydroxy, carboxy, carboxy C_1 - C_6 alkyl, C_1 - C_6 alkyloxy-carbonyl, aryloxy, aryl C_1 - C_6 alkyloxy, aryl C_1 - C_6 alkyl-oxy C_1 - C_6 alkyl, thio, C_1 - C_6 alkylthio, C_1 - C_6 alkylthio C_1 - C_6 alkyl, arylthio, aryl C_1 - C_6 alkylthio, aryl C_1 - C_6 alkylthio C_1 - C_6 alkyl, NR_7R_8 , C_1 - C_6 alkyl-amino C_1 - C_6 alkyl, aryl C_1 - C_6 alkylamino C_1 - C_6 alkyl, di(aryl C_1 - C_6 alkyl)-amino C_1 - C_6 alkyl, C_1 - C_6 alkylcarbonyl, C_1 - C_6 alkylcarbonyl C_1 - C_6 alkyl, aryl C_1 - C_6 alkylcarbonyl, aryl C_1 - C_6 alkylcarbonyl C_1 - C_6 alkyl, C_1 - C_6 alkyl-carboxy, C_1 - C_6 alkyl-carboxy C_1 - C_6 alkyl, arylcarboxy, aryl C_1 - C_6 alkyl-carboxy, aryl C_1 - C_6 alkylcarboxy C_1 - C_6 alkyl, C_1 - C_6 alkylcarbonylamino, C_1 - C_6 alkylcarbonyl-amino C_1 - C_6 alkyl, -carbonyl NR_7C_1 - C_6 alkyl COR_{11} , aryl C_1 - C_6 alkyl-carbonylamino, aryl C_1 - C_6 alkylcarbonylamino C_1 - C_6 alkyl, $CONR_7R_8$, or C_1 - C_6 alkyl $CONR_7R_8$ wherein the alkyl and aryl groups are optionally substituted and R_{11} is NR_7R_8 , or C_1 - C_6 alkyl NR_7R_8 ; or R_3 is



wherein R_{12} , R_{13} , and R_{14} are independently hydrogen, C_1 - C_6 alkyl, aryl, aryl C_1 - C_6 alkyl and the alkyl and aryl groups are optionally substituted;

5 R_4 is hydrogen, hydroxy, C_1 - C_6 alkyl, aryl, aryl C_1 - C_6 alkyl, NR_7R_8 , C_1 - C_6 alkyloxy; wherein the alkyl and aryl groups are optionally substituted;

R_5 is hydroxy, C_1 - C_6 alkyl, aryl, aryl C_1 - C_6 alkyl, CF_3 , NR_7R_8 ; wherein the alkyl and aryl groups are optionally substituted;

10 R_6 is hydrogen, C_1 - C_6 alkyl, aryl, aryl C_1 - C_6 alkyl; wherein the alkyl and aryl groups are optionally substituted;

R_7 and R_8 are independently selected from hydrogen, C_1 - C_6 alkyl, aryl, aryl C_1 - C_6 alkyl, C_1 - C_6 alkyl-carbonyl, arylcarbonyl, aryl C_1 - C_6 alkyl-carbonyl, C_1 - C_6 alkyl-carboxy or aryl C_1 - C_6 alkylcarboxy wherein the alkyl and aryl groups are optionally substituted; or
15 R_7 and R_8 are taken together with the nitrogen to which they are attached forming a cyclic or bicyclic system containing 3 to 11 carbon atoms and 0 to 2 additional heteroatoms selected from nitrogen, oxygen or sulfur, the ring system can optionally be substituted with at least one C_1 - C_6 alkyl, aryl, aryl C_1 - C_6 alkyl, hydroxy, C_1 - C_6 alkyloxy, aryl C_1 - C_6 alkyloxy, C_1 - C_6 alkyloxy C_1 - C_6 alkyl, NR_9R_{10} or C_1 - C_6 alkylamino- C_1 - C_6 alkyl,
20 wherein R_9 and R_{10} are independently selected from hydrogen, C_1 - C_6 alkyl, aryl, aryl C_1 - C_6 alkyl, C_1 - C_6 alkylcarbonyl, arylcarbonyl, aryl C_1 - C_6 alkylcarbonyl, C_1 - C_6 alkylcarboxy or aryl C_1 - C_6 alkylcarboxy; wherein the alkyl and aryl groups are optionally substituted; or

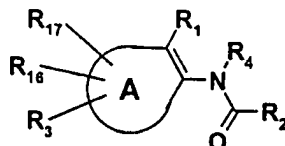
25 R_7 and R_8 are independently a saturated or partial saturated cyclic 5, 6 or 7 membered amine or lactam;

or a salt thereof with a pharmaceutically acceptable acid or base, or any optical isomer or mixture of optical isomers, including a racemic mixture, or any tautomeric
30 forms.

Further, preferred compounds of the invention are compounds of formula 1a or formula 1c wherein R_{16} and R_{17} are hydrogen.

The invention will in its broadest aspect cover the following compounds:

5 of Formula 1b:



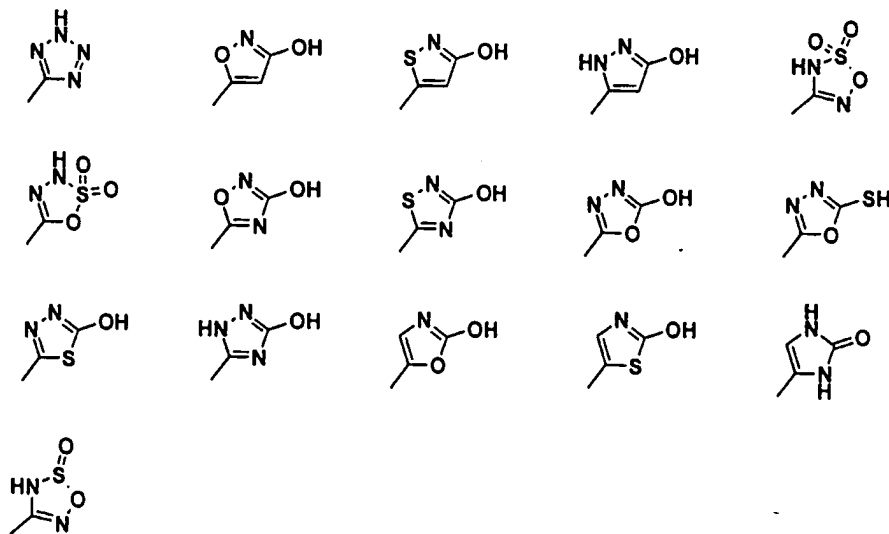
Formula 1b

10

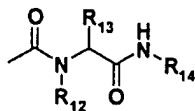
wherein

A is together with the double bond in Formula 1b is aryl;

R_1 is COR_5 , OR_6 , CF_3 , nitro, cyano, SO_3H , $\text{SO}_2\text{NR}_7\text{R}_8$, $\text{PO}(\text{OH})_2$, $\text{CH}_2\text{PO}(\text{OH})_2$,
 15 $\text{CHFPO}(\text{OH})_2$, $\text{CF}_2\text{PO}(\text{OH})_2$, $\text{C}(\text{=NH})\text{NH}_2$, NR_7R_8 ; or
 selected from the following 5-membered heterocycles:



or R_1 is



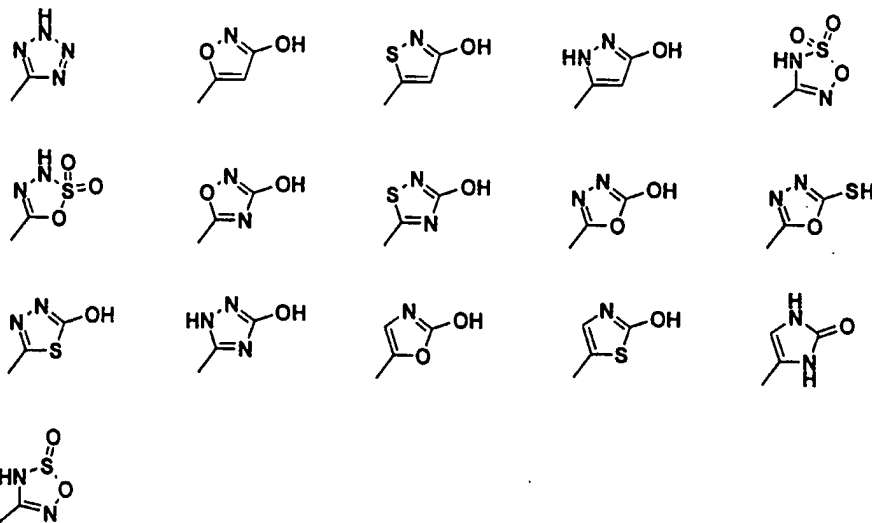
5

wherein R_{12} , R_{13} , and R_{14} are independently hydrogen, C_1 - C_6 alkyl, aryl, aryl C_1 - C_6 alkyl and the alkyl and aryl groups are optionally substituted;

R_2 is COR_5 , OR_6 , CF_3 , nitro, cyano, SO_3H , $SO_2NR_7R_8$, $PO(OH)_2$, $CH_2PO(OH)_2$,

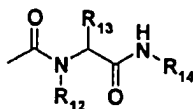
10 $CHFPO(OH)_2$, $CF_2PO(OH)_2$, $C(=NH)NH_2$, NR_7R_8 ; or

selected from the following 5-membered heterocycles:



R_3 , R_{16} and R_{17} are independently hydrogen, halo, nitro, cyano, trihalomethyl, C_1 -
 15 C_6 alkyl, aryl, aryl C_1 - C_6 alkyl, hydroxy, carboxy, carboxy C_1 - C_6 alkyl, C_1 -
 C_6 alkyloxycarbonyl, aryloxycarbonyl, aryl C_1 - C_6 alkyloxycarbonyl, C_1 - C_6 alkyloxy, C_1 -
 C_6 alkyloxy C_1 - C_6 alkyl, aryloxy, aryl C_1 - C_6 alkyloxy, aryl C_1 - C_6 alkyloxy C_1 - C_6 alkyl, thio, C_1 -
 C_6 alkylthio, C_1 - C_6 alkylthio C_1 - C_6 alkyl, arylthio, aryl C_1 - C_6 alkylthio, aryl C_1 - C_6 alkylthio C_1 -
 C_6 alkyl, NR_7R_8 , C_1 - C_6 alkylamino C_1 - C_6 alkyl, aryl C_1 - C_6 alkylamino C_1 - C_6 alkyl, di(aryl C_1 -
 20 C_6 alkyl)amino C_1 - C_6 alkyl, C_1 - C_6 alkylcarbonyl, C_1 - C_6 alkylcarbonyl- C_1 - C_6 alkyl, aryl C_1 -
 C_6 alkylcarbonyl, aryl C_1 - C_6 alkylcarbonyl C_1 - C_6 alkyl, C_1 - C_6 alkylcarboxy, C_1 -
 C_6 alkylcarboxy C_1 - C_6 alkyl, arylcarboxy, arylcarboxy C_1 - C_6 alkyl, aryl C_1 - C_6 alkylcarboxy,

- arylC₁-C₆alkylcarboxyC₁-C₆alkyl, C₁-C₆alkylcarbonylamino, C₁-C₆alkylcarbonylaminoC₁-C₆alkyl, -carbonylNR₇C₁-C₆alkylCOR₁₁, arylC₁-C₆alkylcarbonylamino, arylC₁-C₆alkylcarbonylaminoC₁-C₆alkyl, CONR₇R₈, or C₁-C₆alkylCONR₇R₈ wherein the alkyl and aryl groups are optionally substituted and R₁₁ is NR₇R₈, or C₁-C₆alkylNR₇R₈; or R₃ is



- wherein R₁₂, R₁₃, and R₁₄ are independently hydrogen, C₁-C₆alkyl, aryl, arylC₁-C₆alkyl and the alkyl and aryl groups are optionally substituted;

R₄ is hydrogen, hydroxy, C₁-C₆alkyl, aryl, arylC₁-C₆alkyl, NR₇R₈, C₁-C₆alkyloxy; wherein the alkyl and aryl groups are optionally substituted;

R₅ is hydroxy, C₁-C₆alkyl, aryl, arylC₁-C₆alkyl, CF₃, NR₇R₈; wherein the alkyl and aryl groups are optionally substituted;

R₆ is hydrogen, C₁-C₆alkyl, aryl, arylC₁-C₆alkyl; wherein the alkyl and aryl groups are optionally substituted;

R₇ and R₈ are taken together with the nitrogen to which they are attached forming a saturated, partially saturated or aromatic cyclic, bicyclic or tricyclic ring system containing 3 to 14 carbon atoms and 0 to 3 additional heteroatoms selected from nitrogen, oxygen or sulfur, the ring system can optionally be substituted with at least one C₁-C₆alkyl, aryl, arylC₁-C₆alkyl, hydroxy, oxo, C₁-C₆alkyloxy, arylC₁-C₆alkyloxy, C₁-C₆alkyloxyC₁-C₆alkyl, NR₉R₁₀ or C₁-C₆alkylaminoC₁-C₆alkyl, wherein R₉ and R₁₀ are independently selected from hydrogen, C₁-C₆alkyl, aryl, arylC₁-C₆alkyl, C₁-C₆alkylcarbonyl, arylcarbonyl, arylC₁-C₆alkylcarbonyl, C₁-C₆alkylcarboxy or arylC₁-C₆alkylcarboxy; wherein the alkyl and aryl groups are optionally substituted; or R₇ and R₈ are independently a saturated or partial saturated cyclic 5, 6 or 7 membered amine, imide or lactam;

or a salt thereof with a pharmaceutically acceptable acid or base, or any optical isomer or mixture of optical isomers, including a racemic mixture, or any tautomeric forms.

5

The following compounds are preferred:

- 2-(Oxalyl-amino)benzoic acid;
- 2-Methyl-6-(oxalyl-amino)benzoic acid;
- 10 2-(Oxalyl-amino)-5-iodo-benzoic acid;
- 2-(Aminooxalyl-amino)benzoic acid;
- 2-(Oxalyl-amino)-4-fluoro-benzoic acid;
- 2-(Oxalyl-amino)-4-(3-hydroxy-pyridin-2-ylsulfanyl)benzoic acid;
- 2-(Oxalyl-amino)-4-(1-imidazol-2-ylsulfanyl)benzoic acid;
- 15 2-(Oxalyl-amino)-4-(benzooxazol-2-ylsulfanyl)benzoic acid;
- 2-(Oxalyl-amino)-4-(4-phenyl-thiazol-2-ylsulfanyl)benzoic acid;
- 2-(Oxalyl-amino)-4-(3-ethoxycarbonylphenoxy)benzoic acid;
- 2-(Oxalyl-amino)-4-benzylamino-benzoic acid;
- 2-(Oxalyl-amino)-4-(cyclopropylmethyl-amino)benzoic acid;
- 20 2-(Oxalyl-amino)-4-(3-methyl-butylamino)benzoic acid;
- 2-(Hydrazinooxalyl-amino)benzoic acid;
- 3-(Oxalyl-amino)naphthalene-2-carboxylic acid;
- 2-((5-Thioxo-4,5-dihydro[1,3,4]oxadiazole-2-carbonyl)-amino)benzoic acid;
- N-(2-(3,5-Dichloro-phenoxy)phenyl)oxalamic acid;
- 25 2-(Oxalyl-amino)-3-hydroxy-benzoic acid;
- 2-(Oxalyl-amino)-5-hydroxy-benzoic acid;
- 2-(Oxalyl-amino)-4-nitro-benzoic acid;
- 2-(Oxalyl-amino)-3,5-diiodo-benzoic acid;
- 2-(Oxalyl-amino)-3,5-dimethyl-benzoic acid;
- 30 2-(Oxalyl-amino)-5-methyl-benzoic acid;
- 2-(Oxalyl-amino)-4,5-dimethoxy-benzoic acid;
- 3-(Oxalyl-amino)-phthalic acid;

- 2-(Oxalyl-amino)-4-(2-hydroxy-cyclohexylamino)benzoic acid;
2-(Oxalyl-amino)-4-((2-hydroxy-ethyl)-methyl-amino)benzoic acid;
2-(Oxalyl-amino)-4-(2,2,6,6-tetramethyl-piperidin-4-ylamino)benzoic acid;
2-(2-Hydroxy-cyclohexylamino)-6-(oxalyl-amino)benzoic acid;
5 2-[(2-Hydroxy-ethyl)-methyl-amino]-6-(oxalyl-amino)benzoic acid;
2-(Oxalyl-amino)-6-(2,2,6,6-tetramethyl-piperidin-4-ylamino)benzoic acid;
2-(Oxalyl-amino)-4-(pyridin-3-ylamino)benzoic acid;
2-(Oxalyl-amino)-4-(3-hydroxy-piperidin-1-yl)benzoic acid;
2-(Oxalyl-amino)-4-(1-methyl-1H-imidazol-2-ylsulfanyl)benzoic acid;
10 2-(Oxalyl-amino)-6-(pyridin-3-ylamino)benzoic acid;
2-(3-Hydroxy-piperidin-1-yl)-6-(oxalyl-amino)benzoic acid;
2-(1-Methyl-1H-imidazol-2-ylsulfanyl)-6-(oxalyl-amino)benzoic acid;
2-(Oxalyl-amino)-4-(2-morpholin-4-yl-ethylamino)benzoic acid;
2-(Oxalyl-amino)-4-(benzenesulfonyl-(2-morpholin-4-yl-ethyl)-amino)benzoic acid;
15 2-(Oxalyl-amino)-4-[acetyl-(2-morpholin-4-yl-ethyl)-amino]benzoic acid;
2-(2-Morpholin-4-yl-ethylamino)-6-(oxalyl-amino)benzoic acid;
2-(Benzenesulfonyl-(2-morpholin-4-yl-ethyl)-amino)-6-(oxalyl-amino)benzoic acid;
2-(Acetyl-(2-morpholin-4-yl-ethyl)-amino)-6-(oxalyl-amino)benzoic acid;
2-(Oxalyl-amino)-4-cyclopentylamino-benzoic acid;
20 2-(Oxalyl-amino)-4-(benzenesulfonyl-cyclopentyl-amino)benzoic acid;
2-(Oxalyl-amino)-4-(acetyl-cyclopentyl-amino)benzoic acid;
2-Cyclopentylamino-6-(oxalyl-amino)benzoic acid;
2-(Benzenesulfonyl-cyclopentyl-amino)-6-(oxalyl-amino)benzoic acid;
2-(Acetyl-cyclopentyl-amino)-6-(oxalyl-amino)benzoic acid;
25 4-(Oxalyl-amino)-biphenyl-3-carboxylic acid;
3-(Oxalyl-amino)-isonicotinic acid;
5-(Oxalyl-amino)-2,6-dioxo-1,2,3,6-tetrahydro-pyrimidine-4-carboxylic acid;
2-(Oxalyl-amino)-nicotinic acid;
6-Chloro-5-isopropylamino-3-(oxalyl-amino)-pyrazine-2-carboxylic acid;
30 5,6-Dichloro-3-(oxalyl-amino)-pyrazine-2-carboxylic acid;
3-(Oxalyl-amino)-pyridazine-4-carboxylic acid;

4-(Oxalyl-amino)-pyrimidine-5-carboxylic acid;

3-(Oxalyl-amino)-pyrazine-2-carboxylic acid;

PHARMACOLOGICAL METHODS

- 5 The compounds are evaluated for biological activity with a truncated form of PTP1B (corresponding to the first 321 amino acids), which was expressed in *E. coli* and purified to apparent homogeneity using published procedures well-known to those skilled in the art. The enzyme reactions are carried out using standard conditions essentially as described by Burke et al. (*Biochemistry* **35**; 15989-15996 (1996)). The
- 10 assay conditions are as follows. Appropriate concentrations of the compounds of the invention are added to the reaction mixtures containing different concentrations of the substrate, *p*-nitrophenyl phosphate (range: 0.16 to 10 mM - final assay concentration). The buffer used was 100 mM sodium acetate pH 5.5, 50 mM sodium chloride, 0.1 % (w/v) bovine serum albumin and 5 mM dithiothreitol (total volume 100 μ l).
- 15 The reaction was started by addition of the enzyme and carried out in microtiter plates at 25° C for 60 minutes. The reactions are stopped by addition of NaOH. The enzyme activity was determined by measurement of the absorbance at 405 nm with appropriate corrections for absorbance at 405 nm of the compounds and *p*-nitrophenyl phosphate. The data are analyzed using nonlinear regression fit to classical Michaelis Menten enzyme kinetic models. Inhibition is expressed as K_i values in
- 20 μ M. The results of representative experiments are shown in Table 1

Table 1

Inhibition of classical PTP1B by compounds
of the invention

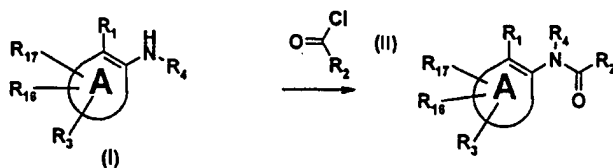
Example no.	PTP1B
	K_i values (μ M)
1	20
2	9.9

Further, the compounds are evaluated for biological activity as regards their effect as inhibitors of PTP α in essentially the same way as described for inhibition of PTP1B. Derived from their activity as evaluated above the compounds of the invention may be useful in the treatment of diseases selected from the group consisting of type I diabetes, type II diabetes, impaired glucose tolerance, insulin resistance and obesity. Furthermore, derived from their activity as evaluated above, the compounds of the invention may be useful in the treatment of diseases selected from the group consisting of immune dysfunctions including autoimmunity, diseases with dysfunctions of the coagulation system, allergic diseases including asthma, osteoporosis, proliferative disorders including cancer and psoriasis, diseases with decreased or increased synthesis or effects of growth hormone, diseases with decreased or increased synthesis of hormones or cytokines that regulate the release of/or response to growth hormone, diseases of the brain including Alzheimer's disease and schizophrenia, and infectious diseases.

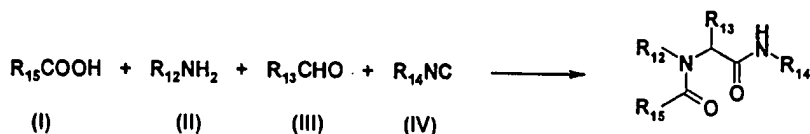
THE SYNTHESIS OF THE COMPOUNDS

In accordance with one aspect of the invention, the compounds of the invention are prepared as illustrated in the following reaction scheme:

Method A

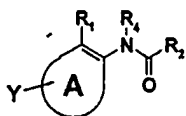


By allowing an amino substituted compound of formula (I) to react with an activated carboxylic acid (e.g. an acid chloride) of formula (II), wherein A, R₁, R₂, R₃, R₄, R₁₆ and R₁₇ are defined as above.

Method B

- 5 By allowing a carboxylic acid (I), a primary amine (II) and an aldehyde (III) to react with a isocyanide (IV) wherein R_{12} , R_{13} , R_{14} , and R_{15} are independently selected from the group consisting of hydrogen, C_1 - C_6 alkyl, aryl, aryl C_1 - C_6 alkyl as defined above and the alkyl and aryl groups are optionally substituted as defined above; or R_{12} , R_{13} , R_{14} , and R_{15} are independently selected from

10



wherein Y indicates attachment point for R_{12} , R_{13} , R_{14} , and R_{15} and A, R_1 , R_2 and R_4 are defined as above.

- 15 In a preferred method, the above described four component Ugi reaction can be carried out by attaching any one of the components to a solid support. Hence, the synthesis can be accomplished in a combinatorial chemistry fashion.

- General procedure for the Preparation of Acetoxymethyl Esters (C.Schultz et al, The
 20 Journal of Biological Chemistry, **1993**, 268, 6316-6322.): A carboxylic acid (1 equivalent) was suspended in dry acetonitrile (2 ml per 0.1 mmol). Diisopropyl amine (3.0 equivalents) was added followed by bromomethyl acetate (1.5 equivalents). The mixture was stirred under nitrogen overnight at room temperature. Acetonitrile was removed under reduced pressure to yield an oil which was diluted in ethylacetate
 25 and washed water (3 x). The organic layer was dried over anhydrous magnesium sulfate. Filtration followed by solvent removal under reduced pressure afforded a crude oil. The product was purified by column chromatography on silica gel, using an appropriate solvent system.

The present invention also has the objective of providing suitable topical, oral, and parenteral pharmaceutical formulations for use in the novel methods of treatment of the present invention. The compounds of the present invention may be administered orally as tablets, aqueous or oily suspensions, lozenges, troches, powders, granules, emulsions, capsules, syrups or elixirs. The composition for oral use may contain one or more agents selected from the group of sweetening agents, flavoring agents, coloring agents and preserving agents in order to produce pharmaceutically elegant and palatable preparations. The tablets contain the acting ingredient in admixture with non-toxic pharmaceutically acceptable excipients which are suitable for the manufacture of tablets. These excipients may be, for example, (1) inert diluents, such as calcium carbonate, lactose, calcium phosphate or sodium phosphate; (2) granulating and disintegrating agents, such as corn starch or alginic acid; (3) binding agents, such as starch, gelatin or acacia; and (4) lubricating agents, such as magnesium stearate, stearic acid or talc. These tablets may be uncoated or coated by known techniques to delay disintegration and absorption in the gastrointestinal tract and thereby provide a sustained action over a longer period. For example, a time delay material such as glyceryl monostearate or glyceryl distearate may be employed. Coating may also be performed using techniques described in the U.S. Patent Nos. 4,256,108; 4,160,452; and 4,265,874 to form osmotic therapeutic tablets for control release.

Formulations for oral use may be in the form of hard gelatin capsules wherein the active ingredient is mixed with an inert solid diluent, for example, calcium carbonate, calcium phosphate or kaolin. They may also be in the form of soft gelatin capsules wherein the active ingredient is mixed with water or an oil medium, such as peanut oil, liquid paraffin or olive oil.

Aqueous suspensions normally contain the active materials in admixture with excipients suitable for the manufacture of aqueous suspension. Such excipients may be (1) suspending agent such as sodium carboxymethyl cellulose, methyl cellulose, hydroxypropylmethyl-cellulose, sodium alginate, polyvinylpyrrolidone, gum tragacanth and gum acacia; (2) dispersing or wetting agents which may be (a) naturally occurring phosphatide such as lecithin; (b) a condensation product of an alkylene oxide

with a fatty acid, for example, polyoxyethylene stearate; (c) a condensation product of ethylene oxide with a long chain aliphatic alcohol, for example, heptadecaethylenoxycetanol; (d) a condensation product of ethylene oxide with a partial ester derived from a fatty acid and hexitol such as polyoxyethylene sorbitol monooleate, or (e) a
5 condensation product of ethylene oxide with a partial ester derived from fatty acids and hexitol anhydrides, for example polyoxyethylene sorbitan monooleate.

The pharmaceutical compositions may be in the form of a sterile injectable aqueous or oleagenous suspension. This suspension may be formulated according to known methods using those suitable dispersing or wetting agents and suspending agents
10 which have been mentioned above. The sterile injectable preparation may also be a sterile injectable solution or suspension in a non-toxic parenterally-acceptable diluent or solvent, for example, as a solution in 1,3-butanediol. Among the acceptable vehicles and solvents that may be employed are water, Ringer's solution, and isotonic sodium chloride solution. In addition, sterile, fixed oils are conventionally employed as a solvent or suspending medium. For this purpose, any bland fixed oil may be employed
15 including synthetic mono- or diglycerides. In addition, fatty acids such as oleic acid find use in the preparation of injectables.

The Compounds of the invention may also be administered in the form of suppositories for rectal administration. These compositions can be prepared by mixing the
20 drug with a suitable non-irritating excipient which is solid at ordinary temperature but liquid at the rectal temperature and will therefore melt in the rectum to release the drug. Such materials are cocoa butter and polyethylene glycols.

The compounds of the present invention may also be administered in the form of liposome delivery systems, such as small unilamellar vesicles, large unilamellar vesicles, and multilamellar vesicles. Liposomes can be formed from a variety of phospholipids, such as cholesterol, stearylamine, or phosphatidylcholines.
25

For topical use, creams, ointments, jellies, solutions or suspensions, etc., containing the compounds of Formula 1 are employed.

Dosage levels of the compounds of the present invention are of the order of about
30 0.5 mg to about 100 mg per kilogram body weight, with a preferred dosage range between about 20 mg to about 50 mg per kilogram body weight per day (from about

25 mg to about 5 g's per patient per day). The amount of active ingredient that may be combined with the carrier materials to produce a single dosage will vary depending upon the host treated and the particular mode of administration. For example, a formulation intended for oral administration to humans may contain 5 mg to 1 g of an active compound with an appropriate and convenient amount of carrier material which may vary from about 5 to about 95 percent of the total composition. Dosage unit forms will generally contain between from about 5 mg to about 500 mg of active ingredient.

It will be understood, however, that the specific dose level for any particular patient will depend upon a variety of factors including the activity of the specific compound employed, the age, body weight, general health, gender, diet, time of administration, route of administration, rate of excretion, drug combination and the severity of the particular disease undergoing therapy. The dosage needs to be individualized by the clinician.

EXAMPLES

The process for preparing compounds of Formula 1 and preparations containing them is further illustrated in the following examples, which, however, are not to be construed as limiting.

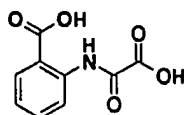
Hereinafter, TLC is thin layer chromatography, CDCl_3 is deuterio chloroform, CD_3OD is tetradeutero methanol and DMSO-d_6 is hexadeutero dimethylsulfoxide. The structures of the compounds are confirmed by either elemental analysis or NMR, where peaks assigned to characteristic protons in the title compounds are presented where appropriate. ^1H NMR shifts (δ_{H}) are given in parts per million (ppm) downfield from tetramethylsilane as internal reference standard. M.p.: is melting point and is given in $^{\circ}\text{C}$ and is not corrected. Column chromatography was carried out using the technique described by W.C. Still *et al.*, *J. Org. Chem.* 43: 2923 (1978) on Merck silica gel 60 (Art. 9385). HPLC analyses are performed using $5\mu\text{m}$ C18 4 x 250 mm

column eluted with various mixtures of water and acetonitrile, flow = 1 ml/min, as described in the experimental section.

Compounds used as starting material are either known compounds or compounds which can readily be prepared by methods known per se.

5

EXAMPLE 1



2-(Oxalyl-amino)benzoic acid:

10

To a stirred solution of anthranilic acid (20.1 g, 0.15 mol) in dry tetrahydrofuran (250 ml) was added dropwise ethyl oxalyl chloride (10.0 g, 0.073 mol). The resulting reaction mixture was stirred at room temperature for 15 min. filtered and the solvent evaporated in vacuo affording crude 16.4 g (94 %) of 2-(ethoxyoxalyl-amino)benzoic acid as an oil.

15

To a solution of the above benzoic acid (10.0 g, 42 mmol) in ethanol (350 ml) was added a solution of sodium hydroxide (3.7 g, 92 mmol) in water (100 ml). The resulting reaction mixture was stirred at room temperature for 60 h. Concentrated hydrochloric acid was added to pH = 1 and the precipitate was filtered off and washed with water (3 x 100 ml), diethyl ether (3 x 80 ml) and dried in vacuo affording 7.1 g (81 %) of the title compound as a solid.

20

M.p.: 214 - 215 °C:

25

Calculated for $C_9H_7NO_5$, 0.2 H_2O ;

C, 51.68 %; H, 3.37 %; N, 6.70 %. Found:

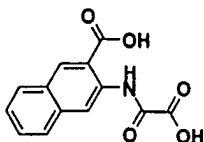
C, 50.96 %; H, 3.32 %; N, 6.52 %.

30

By a similar procedure as described in Example 1 the following compounds have been prepared.

EXAMPLE 2

5



3-(Oxalyl-amino)naphthalene-2-carboxylic acid:

M.p.: 227 - 228 °C:

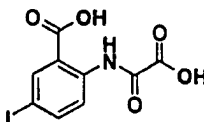
10 Calculated for $C_{13}H_9NO_5$:

C, 60.24 %; H, 3.50 %; N, 5.40 %. Found:

C, 59.98 %; H, 3.46 %; N, 5.25 %.

15

EXAMPLE 3



2-(Oxalyl-amino)-5-iodo-benzoic acid:

20 MS(ES): $m/z = 326$ (M+1)

Calculated for $C_9H_6NO_5$, $0.75 \times H_2O$:

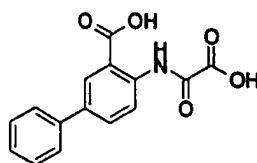
C, 31.01 %; H, 2.17 %; N, 4.02 %. Found:

C, 31.14 %; H, 2.33 %; N, 3.76 %.

25

EXAMPLE 4

52



4-(Oxalyl-amino)-biphenyl-3-carboxylic acid:

To a suspension of 5-bromo-2-amino-benzoic acid methyl ester (3.0 g, 13.04 mmol), tetrakis(triphenylphosphine)palladium(0) (0.5 g, 0.44 mmol), toluene (40 ml) and 2 N aqueous sodium carbonate (14.8 ml) was added a solution of phenylboronic acid (2.2 g, 17.73 mmol) in methanol (10 ml) at room temperature. The resulting reaction mixture was heated at reflux temperature for 4 h. cooled and diluted with water (50 ml). The insoluble matter was filtered off and the phases were separated. The aqueous phase was extracted with ethyl acetate (100 ml) and the combined organic phases were washed with water (2 x 80 ml), diluted aqueous ammoniac (80 ml) and saturated aqueous sodium chloride (80 ml). The organic phase was dried (MgSO₄), filtered and evaporated in vacuo affording 3.4 g of crude 4-amino-biphenyl-3-carboxylic acid methyl ester which was purified on silicagel (1 l) using a mixture of ethyl acetate and heptane (1:3) as eluent. Pure fractions were collected and evaporated in vacuo affording 2.7 g (91 %) of 4-amino-biphenyl-3-carboxylic acid methyl ester.

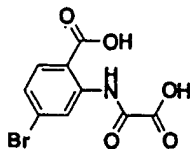
4-Amino-biphenyl-3-carboxylic acid methyl ester was converted into the title compound by a similar procedure as described in Example 1.

M.p.: 223 - 224 °C.

Calculated for C₁₅H₁₁NO₅, 0.5 x H₂O;

C, 61.23 %; H, 4.11 %; N, 4.76 %. Found:

C, 60.96 %; H, 4.01 %; N, 4.62 %.

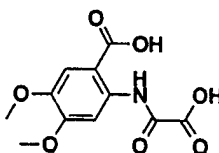
EXAMPLE 5

5 4-Bromo-2-(oxalyl-amino)-benzoic acid:

^1H NMR (400 MHz, CD_3OD) δ 8.71 (d, $J = 7.5$ Hz, 1H), 8.25 (s, 1H), 7.80 (d, $J = 7.5$ Hz, 1H).

MS: ESI (-): 288 [$\text{M}-1(^{81}\text{Br})$], 287 [$\text{M}-1(^{80}\text{Br})$].

10

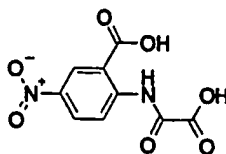
EXAMPLE 6

15 4,5-Dimethoxy-2-(oxalyl-amino)-benzoic acid:

^1H NMR (400 MHz, CD_3OD) δ 8.42 (s, 1H), 7.60 (s, 1H), 3.95 (s, 3H), 3.86 (s, 3H).

MS: ESI (-): 268 [$\text{M}-1$].

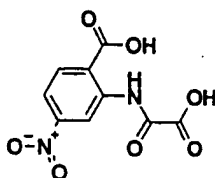
20

EXAMPLE 7

25 5-Nitro-2-(oxalyl-amino)-benzoic acid:

^1H NMR (400 MHz, CD_3OD) δ 8.90 (d, $J = 7.5\text{Hz}$, 2H), 8.42 (s, 1H).

MS: ESI (-): 253 [$\text{M}-1$].

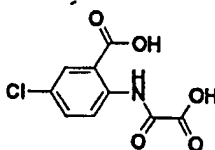
EXAMPLE 84-Nitro-2-(oxalyl-amino)-benzoic acid:

5

¹H NMR (400 MHz, CD₃OD) δ 9.60 (s, 1H), 8.36 (m, 1H), 8.02 (m, 1H).

MS: ESI (-): 253 [M-1].

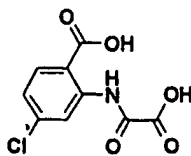
10

EXAMPLE 95-Chloro-2-(oxalyl-amino)-benzoic acid:

15

¹H NMR (400 MHz, CD₃OD) δ 8.72 (d, *J* = 7.5 Hz, 1H), 8.10 (s, 1H), 7.60 (d, *J* = 7.5 Hz, 1H).

MS: ESI (-): 242 [M-1(³⁵Cl)], 244 [M-1(³⁷Cl)].

EXAMPLE 10

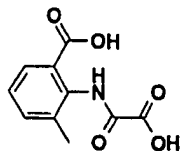
20

4-Chloro-2-(oxalyl-amino)-benzoic acid:

¹H NMR (400 MHz, CD₃OD) δ 8.80 (s, 1H), 8.10 (d, *J* = 7.5 Hz, 1H), 7.22 (d, *J* = 7.5 Hz, 1H).

25

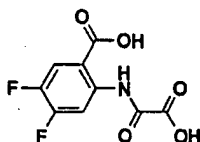
MS: ESI (-): 242 [M-1(³⁵Cl)], 244 [M-1(³⁷Cl)].

EXAMPLE 11

5 3-Methyl-2-(oxalyl-amino)-benzoic acid:

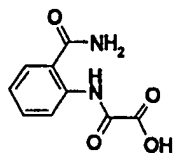
¹H NMR(400 MHz, DMSO-d₆) δ 2.2 (s, 3H), 7.2 - 7.7 (m, 3H), 10.5 (s, 1H), 12.9 (s, 1H).

10

EXAMPLE 12

4,5-Difluoro-2-(oxalyl-amino)-benzoic acid:

15 ¹H NMR (300 MHz, DMSO-d₆) δ 8.03 (m, 1H), 8.61 (dd, 1H), 12.55 (s, 1H, NHCO).

EXAMPLE 13

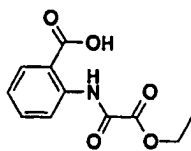
20 N-(2-Carbamoyl-phenyl)-oxalamic acid:

¹H NMR (300 MHz, DMSO-d₆) δ 7.20 (t, 1H), 7.55 (t, 1H), 7.73 (bs, 1H, CONH₂), 7.83 (d, 1H), 8.30 (bs, 1H, CONH₂), 8.52 (d, 1H), 12.9 (s, 1H, NHCO).

25

EXAMPLE 14

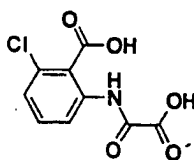
56



2-(Ethoxyoxalyl-amino)-benzoic acid:

¹H NMR (300 MHz, DMSO-d₆) δ 1.33 (t, 3H), 4.30 (q, 2H), 7.24 (t, 1H), 7.65 (t, 1H),
5 8.03 (d, 1H), 8.56 (d, 1H), 12.6 (s, 1H, NHCO).

EXAMPLE 15

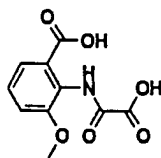


10 6-Chloro-2-(oxalyl-amino)-benzoic acid:

¹H NMR (400 MHz, DMSO-d₆) δ 10.68 (bs, 1H), 8.06 (d, J = 9 Hz, 1H), 7.43 (t, J = 9
Hz, 1H), 7.27 (d, J = 9 Hz, 1H).

15

EXAMPLE 16

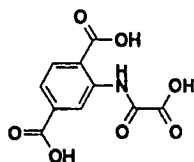


3-Methoxy-2-(oxalyl-amino)-benzoic acid:

20 ¹H NMR (400 MHz, DMSO-d₆) δ 9.98 (bs, 1H), 7.37 - 7.25 (m, 3H), 3.80 (s, 3H).

EXAMPLE 17

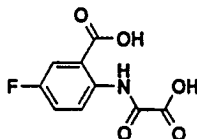
57



2-(Oxalyl-amino)-terephthalic acid:

¹H NMR (400 MHz, DMSO-d₆) δ 7.28 (s, 1H), 8.22 (d, J = 9 Hz, 1H), 7.75 (d, J = 9
5 Hz, 1H).

EXAMPLE 18



5-Fluoro-2-(oxalyl-amino)-benzoic acid:

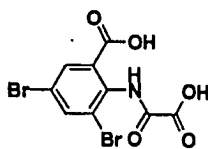
10

¹H NMR (400 MHz, CD₃OD) δ 7.50 (m, 1H), 7.25 (m, 1H), 7.22 (m, 1H).

MS m/z 227.2 (M-1).

15

EXAMPLE 19



3,5-Dibromo-2-(oxalyl-amino)-benzoic acid:

20

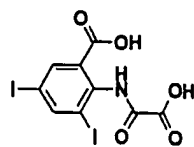
¹H NMR (400 MHz, CD₃OD) δ 8.08 (s, 1H), 8.05 (s, 1H).

MS m/z 366.1 (M-1).

25

EXAMPLE 20

58

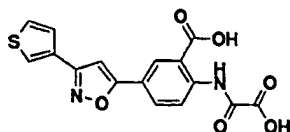


3,5-Diiodo-2-(oxalyl-amino)-benzoic acid:

^1H NMR (400 MHz, CD_3OD) δ 8.45 (s, 1H), 8.25 (s, 1H).

5 MS m/z 460.1 (M-1).

EXAMPLE 21



10

2-(Oxalyl-amino)-5-(3-thiophen-3-yl-isoxazol-5-yl)-benzoic acid:

To a solution of thiophene-3-carboxaldehyde (2.0 g, 18 mmol) in 1,4-dioxane (6.0 ml) was added hydroxylamine hydrochloride (1.24 g, 18 mmol) and triethylamine (2.5 ml, 18 mmol). The mixture was sonicated for 0.5 h and stirred at room temperature for 116 h and at 35 °C for 48 hour. The solvent was removed in vacuo and the residue was dissolved in dichloromethane and washed with water, dried (MgSO_4), filtered and evaporated in vacuo affording 1.97 g (87 %) of thiophene-3-carbaldehyde oxime as an oil.

20 To a solution of the above thiophene-3-carbaldehyde oxime (120 mg, 0.99 mmol) and 2-(*tert*-butoxyoxalyl-amino)-5-ethynyl-benzoic acid methyl ester (100 mg, 0.33 mmol) in tetrahydrofuran (2.5 ml) stirred at room temperature was added 0.75 M bleach (1.3 ml, 0.99 mmol). The solution was first stirred at room temperature for 24 h and then at 35 °C for 24 h. The solvent was evaporated in vacuo and the residue was dissolved in dichloromethane, washed with water, brine and dried (MgSO_4), filtered and evaporated in vacuo. The residual film was purified by preparative TLC affording 21 mg, (15 %) of 2-(*tert*-butoxyoxalyl-amino)-5-(3-thiophen-3-yl-isoxazol-5-yl)-benzoic acid methyl ester as an oil.

¹H NMR (400 MHz, CDCl₃) δ 12.75 (s, 1H), 8.92 (d, 1H, J = 11 Hz), 8.59 (s, 1H), 8.06 (d, 1H, J = 11 Hz), 7.91 (s, 1H), 7.59 (d, 1H, J = 7 Hz), 7.28 (d, 1H, J = 7 Hz), 6.90 (s, 1H), 4.07 (s 3H), 1.65 (s, 9H).

5 The above 2-(*tert*-butoxyoxalyl-amino)-5-(3-thiophen-3-yl-isoxazol-5-yl)-benzoic acid methyl ester (10 mg, 0.023 mmol) was dissolved in 20% trifluoroacetic acid/dichloromethane (0.3 ml) and stirred at room temperature for 21 h. The solvent was removed in vacuo affording 8.4 mg (98 %) of 2-(oxalyl-amino)-5-(3-thiophen-3-yl-isoxazol-5-yl)-benzoic acid methyl ester as a solid.

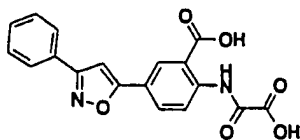
10 ¹H NMR (400 MHz, CD₃OD) δ 12.75 (s, 1H), 8.95 (d, 1H, J = 11 Hz), 8.62 (s, 1H), 8.18 (d, 1H, J = 11 Hz), 7.75 (m, 1H), 7.65 (m, 1H), 7.60 (s, 1H), 4.07 (s, 3H).

To a solution of 2-(oxalyl-amino)-5-(3-thiophen-3-yl-isoxazol-5-yl)-benzoic acid methyl ester (8.4 mg, 0.023 mmol) in methanol (1.5 ml) and tetrahydrofuran (0.5 ml) at room temperature was added 1 M lithium hydroxide (90 μl, 0.090 mmol). The solution was stirred for 48 h. The solvent was removed in vacuo and the residue was redissolved in water. The solution was acidified with 1 N hydrochloric acid to pH = 1 and extracted with ethyl acetate. The combined
15 extracts were washed with brine, dried (MgSO₄), filtered and the solvent evaporated in vacuo affording 6.4 mg (79 %) of the title compound as a solid.
20 ¹H NMR (400 MHz, CD₃OD) δ 8.75 (d, 1H, J=11 Hz), 8.70 (s, 1H), 7.95 (s, 1H), 7.82 (d, 1H, J = 11 Hz), 7.70 (m, 1H), 7.60 (m, 1H).

25 MS m/z: 357(M-1).

EXAMPLE 22

60



2-(Oxalyl-amino)-5-(3-phenyl-isoxazol-5-yl)-benzoic acid:

To a solution of benzaldehyde (2.0 g, 19 mmol) in 1,4-dioxane (6.0 ml) was added
5 hydroxylamine hydrochloride (1.3 g, 19 mmol) and triethylamine (2.6 ml, 19 mmol).
The mixture was sonicated for 0.5 h and stirred at room temperature for 116 h and at
35 °C for 24 h. The solvent was removed in vacuo and the residue was dissolved in
dichloromethane and washed with water, dried (MgSO₄), filtered and evaporated in
vacuo affording 1.9 g (84 %) of benzaldehyde oxime as an oil.

10 ¹H NMR (400 MHz, CDCl₃) δ 8.18 (s, 1H), 7.60 (m, 2H), 7.41 (m, 3H).

To a solution of benzaldehyde oxime (120 mg, 0.99 mmol) and 2-(*tert*-
butoxyoxalyl-amino)-5-ethynyl-benzoic acid methyl ester (100 mg, 0.33 mmol) in
15 tetrahydrofuran (2.5 ml) stirred at room temperature was added 0.75 M bleach
(1.3 ml, 0.99 mmol). The solution was first stirred at room temperature for 24 h
and then at 35 °C for 24 h. The solvent was evaporated in vacuo and the residue
was dissolved in dichloromethane. The solution was washed with water, brine,
dried (MgSO₄), filtered and the solvent evaporated in vacuo. The residue was
20 washed with diethyl ether to get a solid precipitate which was filtered off yielding
59 mg (42 %) of 2-(*tert*-butoxyoxalyl-amino)-5-(3-phenyl-isoxazol-5-yl)-benzoic
acid methyl ester as a solid.

¹H NMR (400 MHz, CDCl₃) δ 12.75 (s, 1H), 8.85 (d, 1H, J = 11 Hz), 8.62 (s, 1H),
8.06 (d, 1H, J = 11 Hz), 7.91 (m, 2H), 7.52 (m, 3H), 6.90 (s, 1H), 4.07 (s 3H),
25 1.65 (s, 9H).

The above 2-(*tert*-butoxyoxalyl-amino)-5-(3-phenyl-isoxazol-5-yl)-benzoic acid
methyl ester (28 mg, 0.07 mmol) was dissolved in 20 % trifluoroacetic
acid/dichloromethane (0.5 ml) and stirred at room temperature for 6 h. The
30 solvent was removed in vacuo affording 25 mg (100 %) of 2-amino-5-(3-phenyl-
isoxazol-5-yl)-benzoic acid methyl ester as a solid.

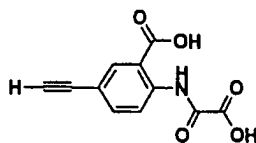
¹H NMR (400 MHz, CDCl₃) δ 12.75 (s, 1H), 8.85 (d, 1H, J = 11 Hz), 8.62 (s, 1H), 8.15 (d, 1H, J = 11 Hz), 7.91 (m, 2H), 7.52 (m, 3H), 6.90 (s, 1H), 4.07 (s 3H)

To a solution of the above 2-amino-5-(3-phenyl-isoxazol-5-yl)-benzoic acid methyl ester (12.5 mg, 0.034 mmol) in methanol (2.5 ml) and tetrahydrofuran (1.0 ml) at room temperature was added 1 M lithium hydroxide (1.4 ml, 0.136 mmol). The solution was stirred for 12 h and the solvent was removed in vacuo. The residue was dissolved in water acidified with 1 N hydrochloric acid to pH = 1 and extracted with ethyl acetate. The organic extract was washed with brine, dried (MgSO₄), filtered and the solvent evaporated in vacuo affording 7.7 mg (64 %) of the title compound as a solid.

¹H NMR (400 MHz, CD₃OD) δ 8.91 (d, 1H, J = 11 Hz), 8.62 (s, 1H), 8.15 (d, 1H, J = 11 Hz), 7.91 (m, 2H), 7.49 (m, 3H), 7.25 (s, 1H), 4.07 (s 3H).

LC/MS m/z: 351(M-1).

EXAMPLE 23



5-Ethynyl-2-(oxalyl-amino)-benzoic acid:

A solution of 2-(*tert*-butoxyoxalyl-amino)-5-iodo-benzoic acid (5.0 g, 12.8 mmol, prepared as described in example 27) and N,N-dimethylformamide di-*tert*butylacetal (12 ml, 51.2 mmol) in toluene (100 ml) was heated at reflux for 20 h. The reaction was cooled to room temperature, concentrated in vacuo and the residue dissolved in ethyl acetate (150 ml). The ethyl acetate phase was washed with water (3 x 35 ml), brine (20 ml) and the volatiles evaporated in vacuo. The residue was purified by silica gel chromatography using 25 % ethyl acetate/hexane as eluent. Pure fractions were combined and concentrated in vacuo to yield 2.3 g of 2-(*tert*-butoxyoxalyl-amino)-5-iodo-benzoic acid *tert*-butyl ester as an oil.

¹H NMR (400 MHz, CDCl₃) δ 8.54 (d, J = 9 Hz, 1H), 8.27 (s, 1H), 7.83 (d, J = 9 Hz, 1H), 1.62 (s, 18H).

5 2-(*tert*-Butoxyoxalyl-amino)-5-iodo-benzoic acid *tert*-butyl ester (0.83 g, 1.86 mmol), trimethylsilyl acetylene (2 ml), and triethylamine (1 ml, 7.44 mmol) were dissolved in N,N-dimethylformamide (5 ml) and the solution purged with argon. Dichlorobis-(triphenylphosphine)palladium(II) (26 mg, 0.15 mmol) and copper(I)iodide (4 mg, 0.15 mmol) were added and the reaction stirred at 60 °C under argon for 5 h. The
10 crude mixture was diluted with ethyl acetate (40 ml) and washed with water (3 x 10 ml) and brine (2 x 10 ml). The solvent was evaporated in vacuo to yield 0.77 g (99 %) of 2-(*tert*-butoxyoxalyl-amino)-5-trimethylsilanylethyny-benzoic acid *tert*-butyl ester. ¹H NMR (400 MHz, CDCl₃) δ 12.59 (s, 1H), 8.71 (d, J = 9 Hz, 1H), 8.07 (d, J = 9 Hz, 1H), 1.62 (s, 9H), 1.61 (s, 9H), 0.25 (s, 9H).

15

2-(*tert*-Butoxyoxalyl-amino)-5-trimethylsilanylethyny-benzoic acid *tert*-butyl ester (0.57 g, 1.37 mmol) was dissolved in tetrahydrofuran (5 ml) and treated with a 0.9 M solution of tetrabutylammonium fluoride and acetic acid (2:3) in tetrahydrofuran (1.7
20 ml, 1.51 mmol) for 3 h. The volatiles were evaporated in vacuo and the crude material extracted into ethyl acetate (35 ml). The ethyl acetate extract was washed with 1N hydrochloric acid (5 ml), saturated sodium bicarbonate (5 ml), brine (5 ml) and evaporated in vacuo affording 0.36 g (76 %) of 2-(*tert*-butoxyoxalyl-amino)-5-ethynyl-benzoic acid *tert*-butyl ester as an oil.

25 ¹H NMR (400 MHz, CDCl₃) δ 8.74 (d, J = 10 Hz, 1H), 8.12 (s, 1H), 7.65 (d, J = 10 Hz, 1H), 3.08 (s, 1H), 1.62 (s, 9H), 1.58 (s, 9H).

2-(*tert*-Butoxyoxalyl-amino)-5-ethynyl-benzoic acid *tert*-butyl ester (0.36 g, 1.04 mmol) was treated with 50 % trifluoroacetic acid/dichloromethane (15 ml) at room
30 temperature for 3 h. The reaction mixture was concentrated in vacuo and the residue was washed with water and diethyl ether affording after drying 0.21 g (86 %) of the title compound.

¹H NMR (400 MHz, DMSO-d₆) δ 8.23 (d, J = 10 Hz, 1H), 8.05 (s, 1H), 7.76 (d, J = 10 Hz, 1H), 4.24 (s, 1H).

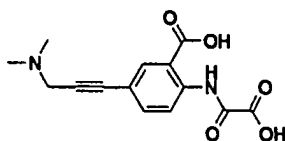
35

LC/MS [M-H]⁻: 232.07

HPLC (254.4 nm): 3.112 s, (49 %).

5

EXAMPLE 24



5-(3-Dimethylamino-prop-1-ynyl)-2-(oxalyl-amino)-benzoic acid:

10 To a solution of 5-iodoanthranilic acid (3.0 g, 11.4 mmol) and N,N-diisopropylethylamine (4 ml, 22.8 mmol) in anhydrous tetrahydrofuran (40 ml) was added imidazol-1-yl-oxo-acetic acid *tert*-butyl ester (4.47 g, 22.8 mmol). The reaction was stirred at room temperature for 3 h. The solvents were evaporated in vacuo and the crude mixture extracted into ethyl acetate (70 ml). The organic extract was
15 washed with 1 % hydrochloric acid (2 x 15 ml) and brine (10 ml) and the solvent was evaporated in vacuo affording 2.8 g (63 %) of 2-(*tert*-butoxyoxalyl-amino)-5-iodo-benzoic acid as a solid.

¹H NMR (400 MHz, CDCl₃) δ 8.57 (d, J = 9 Hz, 1H), 8.43 (d, J = 2 Hz, 1H), 8.00 (dd, J = 9 Hz, 2 Hz, 1H), 1.59 (s, 9H).

20

To a solution of 2-(*tert*-butoxyoxalyl-amino)-5-iodo-benzoic acid (2.1 g, 5.37 mmol) in dichloromethane (15 ml) under nitrogen was added triethylamine (3.75 ml, 26.85 mmol) and N,N-dimethylaminopyridine (0.1 g). Methoxymethyl chloride (1.2 ml, 16.11 mmol) was added and the reaction mixture was stirred for 4 h and concentrated in
25 vacuo to a minimum volume which was loaded directly onto a silica gel column, eluting with 50 % ethyl acetate/hexane. Pure fractions were combined and concentrated to give 1.5 g (64 %) of 2-(*tert*-butoxyoxalyl-amino)-5-iodo-benzoic acid methoxymethyl ester as a solid.

¹H NMR (400 MHz, CDCl₃) δ 8.56 (d, J = 9 Hz, 1H), 8.42 (s, 1H), 7.89 (d, J = 9 Hz, 1H), 5.54 (s, 2H), 3.60 (s, 3H), 1.61 (s, 9H).

30

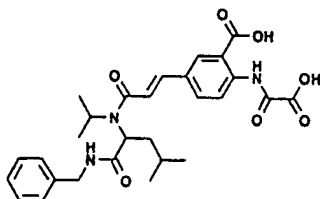
A solution of 2-(*tert*-butoxyoxalyl-amino)-5-iodo-benzoic acid methoxymethyl ester (0.16 g, 0.37 mmol), triethylamine (51 μ l, 0.37 mmol) and 1-dimethylamino-2-propyne (0.12 ml, 1.11 mmol) was prepared in anhydrous acetonitrile (3 ml) and purged with argon. Dichlorobis-(triphenylphosphine)palladium (II) (5 mg, 0.0074 mmol) and copper(I) iodide (1 mg, 0.0074 mmol) were added and the reaction stirred at 60 °C under argon for 18 h. The volatiles were evaporated in vacuo and the residue redissolved in ethyl acetate (10 ml). The organic phase was washed with 1% hydrochloric acid (5 ml) and the aqueous phase extracted with additional ethyl acetate. The combined organic extracts were washed with brine (5 ml), dried (Na_2SO_4) and concentrated to an oil. The crude oil was dissolved in dichloromethane and purified by silica gel chromatography using 5 % methanol/dichloromethane/0.1 % triethylamine as eluent. Pure fractions were combined to give 81 mg (60 %) of 2-(*tert*-butoxyoxalyl-amino)-5-(3-dimethylamino-prop-1-ynyl)-benzoic acid methoxymethyl ester as an oil.

^1H NMR (400 MHz, CDCl_3) δ 12.52 (s, 1H), 8.75 (d, J = 9 Hz, 1H), 8.21 (s, 1H), 7.64 (d, J = 9 Hz, 1H), 5.53 (s, 2H), 3.59 (s, 3H), 3.46 (s, 2H), 2.38 (s, 6H), 1.61 (s, 9H).

2-(*tert*-Butoxyoxalyl-amino)-5-(3-dimethylamino-prop-1-ynyl)-benzoic acid methoxymethyl ester (32.3 mg) was treated with 50 % trifluoroacetic acid/dichloromethane (3 ml) at room temperature for 2 h. The mixture was concentrated to a solid in vacuo and the resulting solid was washed with dichloromethane which afforded 20 mg (83 %) of the title compound as a solid.

^1H NMR (400 MHz, $\text{DMSO}-d_6$) δ 8.60 (d, J = 9 Hz, 1H), 8.05 (s, 1H), 7.60 (d, J = 9 Hz, 1H), 4.07 (s, 2H), 2.73 (s, 6H).

EXAMPLE 25

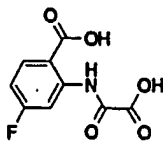


5-(2-(((1-Benzylcarbamoyl-3-methyl-butyl)-isopropyl-carbamoyl)-vinyl)-2-(oxalyl-amino)-benzoic acid:

- 5-(2-(((1-Benzylcarbamoyl-3-methyl-butyl)-isopropyl-carbamoyl)-vinyl)-2-(*tert*-butoxyoxalyl-amino)-benzoic acid methoxymethyl ester (0.14 g, 0.22 mmol, prepared as described in Example 33) was treated with a solution of 50 % trifluoroacetic acid/dichloromethane (6 ml) for 2.5 h. The mixture was concentrated in vacuo and precipitated from water. The resulting crystalline solid was filtered off and dried in vacuo to give 0.10 g (85 %) of the title compound as a solid.
- ¹H NMR (400 MHz, CD₃OD) δ 8.76 (d, J = 9 Hz, 1H), 8.31 (s, 1H), 7.94 (d, J = 9 Hz, 1H), 7.49 (d, J = 16 Hz, 1H), 7.46 - 7.38 (m, 5H), 7.13 (d, J = 16 Hz, 1H), 4.49 (m, 1H), 4.10 (s, 2H), 3.95 (m, 1H), 2.49 (m, 1H), 1.90 (m, 1H), 1.56 (m, 1H), 1.38 (d, J = 6 Hz, 3H), 1.35 (d, J = 7 Hz, 3H), 0.97 (d, J = 6 Hz, 6H).
- LC/MS [M-H]⁻: 522.55.

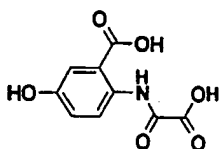
The following compounds were prepared in a similar way as described in Example 1.

EXAMPLE 26



4-Fluoro-2-(oxalyl-amino)-benzoic acid:

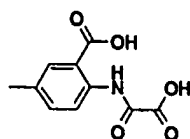
- ¹H NMR (300 MHz, DMSO-d₆) δ 7.11 (m, 1H), 8.12 (m, 1H), 8.42 (dd, 1H), 12.62 (s, 1H, NHCO).

EXAMPLE 275-Hydroxy-2-(oxalyl-amino)-benzoic acid:

5

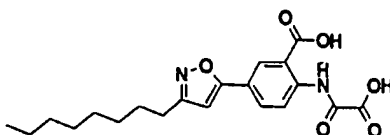
^1H NMR (400 MHz, DMSO- d_6) δ 12.19 (s, 1H) 9.78 (bs, 1H) 8.44 (d, J = 10 Hz, 1H), 7.42 (d, J = 2 Hz, 1H), 7.05 (dd, J = 10 Hz, 2 Hz).

10

EXAMPLE 285-Methyl-2-(oxalyl-amino)-benzoic acid:

15

^1H NMR (400 MHz, DMSO- d_6) δ 12.20 (s, 1H), 8.51 (d, J = 10 Hz, 1H), 7.83 (d, J = 2 Hz, 1H), 7.42 (dd, J = 10 Hz, 2 Hz, 1H), 2.29 (s, 3H).

EXAMPLE 29

20

5-(3-Octyl-isoxazol-5-yl)-2-(oxalyl-amino)-benzoic acid:

To a solution of 2-(*tert*-butoxyoxalyl-amino)-5-iodo-benzoic acid (1.8 g, 4.6 mmol) and potassium carbonate (1.6 g, 11.5 mmol) in acetone (15 ml) was added iodo-methane (3 g). The reaction was heated at reflux for 2 h. after which it was judged complete by TLC analysis. The crude mixture was diluted with ethyl acetate (75 ml),

25

washed with water (2 x 15 ml), and brine (10 ml). The organic phase was concentrated in vacuo to give 1.8 g (94 %) of 2-(*tert*-butoxyoxalyl-amino)-5-iodo-benzoic acid methyl ester.

¹H NMR (400 MHz, CDCl₃) δ 12.52 (s, 1H), 8.53 (d, J = 9 Hz, 1H), 8.39 (s, 1H), 8.87
5 (d, J = 9 Hz, 1H), 3.98 (s, 3H), 1.61 (s, 9H).

2-(*tert*-Butoxyoxalyl-amino)-5-iodo-benzoic acid methyl ester (0.86 g, 2.12 mmol), trimethylsilyl acetylene (2 ml), and triethylamine (1.2 ml, 8.48 mmol) were dissolved in N,N-dimethylformamide (5 ml) and the solution purged with argon. Dichlorobis-
10 (triphenylphosphine)palladium (II) (30 mg, 0.042 mmol) and copper(I)iodide (4 mg, 0.021 mmol) were added and the reaction stirred at 60 °C under argon for 3 h. The crude mixture was diluted with ethyl acetate (40 ml) and washed with water (3 x 10 ml) and brine (2 x 10 ml). The organic phase was evaporated in vacuo to yield 0.8 g (99 %) of 2-(*tert*-Butoxyoxalyl-amino)-5-trimethylsilanylethyny-benzoic acid methyl
15 ester which was used without further purification.

2-(*tert*-Butoxyoxalyl-amino)-5-trimethylsilanylethyny-benzoic acid methyl ester (0.15 g, 0.4 mmol) was dissolved in tetrahydrofuran (2 ml) and treated with a 0.9 M solution of tetrabutylammonium fluoride and acetic acid (2:3) in tetrahydrofuran (0.44 ml, 0.4 mmol) for 3 h. The volatiles were evaporated in vacuo and the crude material extracted into ethyl acetate (25 ml). The ethyl acetate extract was washed with 1N hydrochloric acid (5 ml), saturated sodium bicarbonate (5 ml), brine (5 ml) and evaporated in vacuo affording 0.1 g (83 %) of 2-(*tert*-butoxyoxalyl-amino)-5-ethynyl-benzoic
20 acid methyl ester as an oil.

25 A solution of 2-(*tert*-butoxyoxalyl-amino)-5-ethynyl-benzoic acid methyl ester (0.1 g, 0.33 mmol) and nonaldoxime (0.15 g, 0.99 mmol) in tetrahydrofuran (3 ml) was treated with bleach (0.75 N, 1.3 ml, 0.99 mmol). The reaction was stirred at room temperature for 24 h. TLC analysis showed presence of starting material so the reaction was heated to 35 °C for 12 h. The solvents were removed in vacuo and the crude
30 material was dissolved in ethyl acetate (35 ml), washed with water (2 x 10 ml) and brine (10 ml). The organic extract was evaporated in vacuo affording 0.1 g (66 %) of

2-(*tert*-butoxyoxalyl-amino)-5-(3-octyl-isoxazol-4-yl)-benzoic acid methyl ester as an oil.

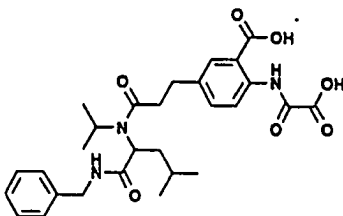
¹H NMR (400 MHz, CDCl₃) δ 8.89 (d, J = 10 Hz, 1H), 8.52 (d, J = 1 Hz, 1H), 7.97 (dd, J = 10, 1 Hz, 1H), 6.41 (s, 1H), 4.02 (s, 3H), 2.73 (t, J = 8 Hz, 2H), 1.70 (m, 2H), 1.62 (s, 9H), 1.37 - 1.25 (bm, 12H), 0.89 (t, J = 8 Hz, 3H).

To a solution of the above (isoxazol-4-yl)-benzoic acid methyl ester (9.1 mg, 0.02 mmol) in 50 % methanol/tetrahydrofuran (2 ml) was added 1N lithium hydroxide (60 μl, 0.06 mmol) and the resulting mixture was stirred at room temperature for 48 h.

The reaction was judged to be incomplete by TLC analysis (30 % methanol/dichloromethane) and additional 1 N lithium hydroxide was added (20 μl, 0.02 mmol). The reaction was stirred for another 72 hours. pH of the reaction mixture was adjusted to around 0 by addition of 1 N hydrochloric acid. The mixture was concentrated *in vacuo* and the crude material was dissolved in ethyl acetate (20 ml). The organic layer was washed with brine (2 x 5 ml) and concentrated *in vacuo* affording 5.4 mg (70 %) of the title compound as a solid.

¹H NMR (400 MHz, CD₃OD) δ 8.87 (d, J = 10 Hz, 1H), 8.55 (s, 1H), 8.04 (d, J = 10 Hz, 1H), 6.74 (s, 1H), 2.70 (t, J = 8 Hz, 2H), 1.72 (m, 2H), 1.38 - 1.20 (bm, 12H), 0.90 (t, J = 8 Hz, 3H).

EXAMPLE 30



5-(2-((1-Benzylcarbamoyl-3-methyl-butyl)-isopropyl-carbamoyl)-ethyl)-2-(oxalyl-amino)-benzoic acid:

To a solution of isopropylamine (0.43 ml, 5.0 mmol) in methanol (5 ml) was added isovaleraldehyde (0.54 ml, 5.0 mmol). After 15 minutes of stirring a solution of benzylisocyanide in tetrahydrofuran (1N, 5 ml, 5.0 mmol) was added followed by acrylic acid (0.34 ml, 5.0 mmol). The reaction was stirred at room temperature for 72 h, the
5 volatiles were removed in vacuo and the resulting oil dissolved in ethyl acetate (40 ml). The organic mixture was washed with 1N hydrochloric acid (10 ml) and brine (10 ml), dried (Na_2SO_4) and the solvent evaporated in vacuo. The crude residue was purified by chromatography using a gradient from 30 % ethyl acetate/hexane to 50 % ethyl acetate/hexane. Pure fraction were collected and the solvent evaporated in
10 vacuo which afforded 1.5 g (100 %) of N-(1-benzylcarbamoyl-3-methyl-butyl)-N-isopropyl-acrylamide as an oil.

^1H NMR (400 MHz, CDCl_3) δ 8.13 (bs, 1H), 7.30 - 7.19 (m, 5H), 6.49 (dd, J = 16 Hz, 12 Hz, 1H), 6.25 (d, J = 16 Hz, 1H), 5.66 (d, J = 12 Hz, 1H), 4.38 (d, J = 6 Hz, 2H), 4.10 - 4.02 (m, 1H), 2.22 - 2.13 (m, 1H), 1.76 - 1.70 (m, 1H), 1.62 - 1.54 (m, 2H),
15 1.25 (d, J = 7 Hz, 3H), 1.20 (d, J = 7 Hz, 3H), 0.94 (d, J = 7 Hz, 3H), 0.90 (d, J = 7 Hz, 3H).

A solution of N-(1-benzylcarbamoyl-3-methyl-butyl)-N-isopropyl-acrylamide (0.55 g, 1.74 mmol), 2-(*tert*-butoxyoxalyl-amino)-5-iodo-benzoic acid methoxymethyl ester
20 (0.5 g, 1.15 mmol), palladium acetate (3.0 mg, 0.023 mmol) and tri(*o*-tolyl)phosphine (10.0 mg, 0.07 mmol) in N,N-dimethylformamide under argon and heated to 100 °C with stirring for 3 h. The reaction was cooled to room temperature and diluted in ethyl acetate (50 ml). The organic phase was washed with water (2 x 15 ml) and brine (10 ml), dried (Na_2SO_4) and evaporated in vacuo. The crude oily material was purified by
25 chromatography using 30 % ethyl acetate/hexane as eluent. Pure fractions were collected and the solvent evaporated in vacuo affording 0.15 g (20 %) of 5-(2-((1-Benzylcarbamoyl-3-methyl-butyl)-isopropyl-carbamoyl)-vinyl)-2-(*tert*-butoxyoxalyl-amino)-benzoic acid methoxymethyl ester as an oil.

^1H NMR (400 MHz, CDCl_3) δ 12.58 (s, 1H), 8.82 (d, J = 9 Hz, 1H), 8.22 (s, 1H), 7.80
30 (d, J = 9 Hz, 1H), 7.60 (d, J = 16 Hz, 1H), 7.30 - 7.22 (m, 5H), 6.80 (d, J = 16 Hz, 1H), 5.58 (s, 2H), 4.43 (bs, 2H), 4.21 - 4.15 (m, 1H), 3.60 (s, 3H), 2.21 - 2.16 (m, 1H),

1.82 - 1.78 (m, 1H), 1.61 (s, 9H), 1.61 - 1.58 (m, 1H), 1.35 (d, J = 7 Hz, 3H), 1.24 (t, J = 7 Hz, 3H), 0.99 (d, J = 7 Hz, 3H), 0.94 (d, J = 7 Hz, 3H).

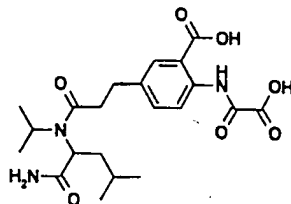
To a solution of 5-(2-((1-Benzylcarbamoyl-3-methyl-butyl)-isopropyl-carbamoyl)-vinyl)-2-(*tert*-butoxyoxalyl-amino)-benzoic acid methoxymethyl ester (10.7 mg, 0.017 mmol) in methanol (1 ml) was added 5 % palladium/carbon (2.2 mg) and the resulting mixture was stirred under hydrogen gas (30 psi) for 3 h. The mixture was filtered through celite and evaporated *in vacuo*. NMR indicated that the reaction was not complete so it was subjected to the hydrogenation conditions for another 4 h. The mixture was filtered and evaporated *in vacuo* again affording 8.9 mg (83 %) of 5-(2-((1-benzylcarbamoyl-3-methyl-butyl)-isopropyl-carbamoyl)-ethyl)-2-(*tert*-butoxyoxalyl-amino)-benzoic acid methoxymethyl ester as an oil.

¹H NMR (400 MHz, CDCl₃) δ 12.41 (s, 1H), 8.68 (d, J = 9 Hz, 1H), 8.07 (bs, 1H), 7.98 (s, 1H), 7.43 (d, J = 9 Hz, 1H), 7.30 - 7.22 (m, 5H), 5.52 (s, 2H), 4.45 - 4.33 (m, 2H), 4.04 - 3.96 (m, 2H), 3.58 (s, 3H), 2.95 (t, J = 7 Hz, 2H), 2.72 - 2.61 (m, 2H), 2.30 (m, 1H), 1.62 (s, 9H), 1.59 (m, 1H partially obscured by neighboring singlet), 1.22 (d, J = 6 Hz, 6H), 0.95 (d, J = 7 Hz, 3H), 0.91 (d, J = 7 Hz, 3H).

5-(2-((1-Benzylcarbamoyl-3-methyl-butyl)-isopropyl-carbamoyl)-ethyl)-2-(*tert*-butoxyoxalyl-amino)-benzoic acid methoxymethyl ester (4 mg, 0.0064 mmol) was dissolved in acetone (3 ml) and treated with 3 drops of 1N hydrochloric acid. The reaction was stirred for 2 days, after which the acetone was evaporated *in vacuo*. The residue was dissolved in ethyl acetate (10 ml), washed with brine (2 x 2ml) and evaporated *in vacuo*. The resulting oil was treated with 20 % trifluoroacetic acid/dichloromethane (3 ml) for 3 h. The volatiles were evaporated *in vacuo* affording 2 mg (61 %) of the title compound as an oil.

¹H NMR (400 MHz, CD₃OD) δ 8.64 (d, J = 9 Hz, 1H), 8.00 (s, 1H), 7.51 (d, J = 9 Hz, 1H), 7.50 - 7.40 (m, 5H), 4.17 (t, J = 8 Hz, 1H), 4.14 (s, 2H), 3.73 (m, 1H), 2.95 (t, J = 6 Hz, 2H), 2.82 - 2.63 (m, 2H), 2.42 (m, 1H), 1.80 (m, 1H), 1.30 (m, 1H), 1.26 (d, J = 6 Hz, 3H), 1.10 (d, J = 6 Hz, 3H), 0.90 (d, J = 6 Hz, 6H).

LC/MS [M-H]⁻: 524.74.

EXAMPLE 31

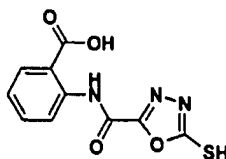
5 5-(2-(((1-Carbamoyl-3-methyl-butyl)-isopropyl-carbamoyl)-ethyl)-2-(oxalyl-amino)-benzoic acid:

5-(2-(((1-Benzylcarbamoyl-3-methyl-butyl)-isopropyl-carbamoyl)-vinyl)-2-(oxalyl-amino)-benzoic acid (33 mg, 0.063 mmol) and 10 % palladium/carbon was mixed in methanol (3 ml) and stirred under hydrogen gas (1 atm) for 18 h. The mixture was
 10 filtered through celite and the volatiles were evaporated in vacuo affording 27 mg (99 %) of the title compound.

¹H NMR (400 MHz, CD₃OD) δ 8.64 (d, J = 9 Hz, 1H), 8.00 (s, 1H), 7.51 (d, J = 9 Hz, 1H), 4.17 (t, J = 8 Hz, 1H), 3.72 (m, 1H), 2.96 (t, J = 6 Hz, 2H), 2.82-2.63 (m, 2H), 2.41 (m, 1H), 1.80 (m, 1H), 1.30 (m, 1H), 1.25 (d, J = 6 Hz, 3H), 1.13 (d, J = 6 Hz,
 15 3H), 0.90 (d, J = 6 Hz, 6H).

LC/MS [M-H]⁻: 435.66

20

EXAMPLE 32

2-((5-Mercapto-[1,3,4]oxadiazole-2-carbonyl)-amino)-benzoic acid:

To a solution of 2-(Ethoxyoxalyl-amino)-benzoic acid (2.0 g, 8.43 mmol) in ethanol (75 ml) was added hydrazine hydrate (0.8 g, 16.86 mmol). The resulting mixture was
 25 stirred at reflux temperature for 3 h. To the cooled reaction was added water (200 ml) and the mixture was acidified with 1 N hydrochloric acid to pH = 4. The precipitate

was filtered off, washed with water and dried in vacuo at 50 °C for 16 h which afforded 1.4 g (69 %) of 2-(hydrazinooxalyl-amino)-benzoic acid as a solid.

To a solution of the above 2-(hydrazinooxalyl-amino)-benzoic acid (1.0 g, 4.15 mmol) in methanol (20 ml) cooled to 0 °C was added potassium hydroxide (0.5 g, 8.72 mmol) and carbondisulfide (0.7 g, 9.54 mmol). The resulting mixture was stirred at reflux temperature for 6 h. To the cooled reaction was added water (100 ml) and the mixture was acidified with 1 N hydrochloric acid to pH = 1. The precipitate was filtered off, washed with water and heptane and dried in vacuo at 50 °C. The dried product (0.65 g) was purified by silica gel (400 ml) chromatography using 5 % acetic acid in ethyl acetate as eluent. Pure fractions were collected and the volatiles were evaporated in vacuo. The residue was washed with water and dried in vacuo at 50 °C for 16 h affording 0.4 g (36 %) of the title compound as a solid.

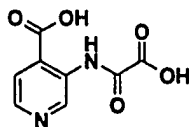
M.p.: 236 - 237 °C

Calculated for $C_{10}H_7N_3O_4S$;

C, 45.28 %; H, 2.66 %; N, 15.84 %. Found:

C, 45.48 %; H, 2.66 %; N, 15.36 %.

EXAMPLE 33



3-(Oxalyl-amino)-isonicotinic acid:

To a stirred solution of 3-amino-isonicotinic acid (0.5 g, 3.62 mmol) and triethylamine (1 ml) in dry tetrahydrofuran (50 ml) at 0 °C was added dropwise ethyl oxalyl chloride (0.5 g, 3.69 mmol). The resulting reaction mixture was stirred at room temperature for 3 h, filtered and the volatiles were evaporated in vacuo. To the residue was

added water (50 ml) and the resulting mixture was extracted with diethyl ether (2 x 50 ml). The organic phase was washed with saturated aqueous sodium chloride (50 ml), dried (MgSO_4), filtered and the solvent evaporated in vacuo affording 0.4 g (46 %) of 3-(ethoxyoxalyl-amino)-isonicotinic acid as a solid.

5 To a solution of the above isonicotinic acid (0.4 g, 1.7 mmol) in ethanol (25 ml) was added a solution of sodium hydroxide (141 mg, 3.53 mmol) in water (10 ml). The resulting reaction mixture was stirred at room temperature for 18 h. The volatiles were evaporated in vacuo and the residue dissolved in water (50 ml) and washed with diethyl ether (50 ml). To the aqueous phase was added 1N hydrochloric acid to pH =
10 1. The precipitate was filtered off and dried in vacuo at 50 °C for 18 h. The dried solid residue was washed with boiling acetone (50 ml) for 5 min. filtered off and dried in vacuo at 50 °C affording 80 mg (22 %) of the title compound as a solid.

M.p.: > 250 °C;

15

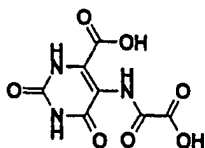
Calculated for $\text{C}_8\text{H}_6\text{N}_2\text{O}_5$;

C, 45.72 %; H, 2.88 %; N, 13.33 %. Found:

C, 45.62 %; H, 2.98 %; N, 13.04 %.

20

EXAMPLE 34



25 5-(Oxalyl-amino)-2,6-dioxo-1,2,3,6-tetrahydro-pyrimidine-4-carboxylic acid:

To a solution of 5-aminoorotic acid (61.1mg, 0.36 mmol) in tetrahydrofuran (1 ml) was added imidazol-1-yl-oxo-acetic acid *tert*-butyl ester (140 mg, 0.71 mmol) and triethylamine (50 μl , 0.36 mmol). The mixture was stirred at room temperature for 20 h. The solvent was removed in vacuo and the residue was dissolved in ethyl acetate

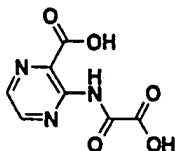
(5.0 ml) and washed with 1 % hydrochloric acid (2 x 2 ml) then water (2 x 2 ml). The organic phase was dried (MgSO_4), filtered and the solvent evaporated in vacuo. The residue was purified by preparative TLC (Kieselgel 60F₂₅₄, 0.5 mm, hexane:ethyl acetate, 80:20) which afforded 30 mg (28 %) of 5-(*tert*-butoxyoxalyl-amino)-2,6-dioxo-1,2,3,6-tetrahydro-pyrimidine-4-carboxylic acid as a solid.

^1H NMR (400 MHz, CDCl_3) δ 1.80 (s, 9H), 7.56 (s, 2H), 8.96 (s, 1H).

5-(*tert*-Butoxyoxalyl-amino)-2,6-dioxo-1,2,3,6-tetrahydro-pyrimidine-4-carboxylic acid (28 mg, 0.094 mmol) was stirred in 20 % trifluoroacetic acid in dichloromethane (1.0 ml) at room temperature for 2 h. The reaction mixture was co-evaporated in vacuo with toluene to complete dryness which afforded 22.6 mg (100 %) of the title compound as a solid.

^1H NMR (400 MHz, CD_3OD) δ 7.30 (s, 2H).

EXAMPLE 35



3-(Oxalyl-amino)-pyrazine-2-carboxylic acid:

To a solution of 3-aminopyrazine-2-carboxylic acid (64.2 mg, 0.46 mmol) in tetrahydrofuran (1 ml) was added imidazol-1-yl-oxo-acetic acid *tert*-butyl ester (181mg, 0.92 mmol) and triethylamine (64.3 μl , 0.46 mmol). The mixture was stirred at room temperature for 20 h. The solvent was removed in vacuo and the residue was dissolved in ethyl acetate (5.0 ml) and washed with 1 % hydrochloric acid (2 x 2 ml) then water

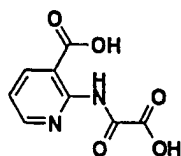
(2 x 2 ml). The organic phase was dried (MgSO₄), filtered and the solvent evaporated in vacuo. The residue was washed with diethyl ether (4 x 1.0 ml) affording 48 mg (39 %) of 3-(*tert*-butoxyoxalyl-amino)-pyrazine-2-carboxylic acid as a solid.

¹H NMR (CDCl₃ + CD₃OD) δ 1.70 (s, 9H), 8.02 (d, 1H, J = 1.5 Hz), 8.36 (d, 1H, J = 1.5 Hz).

3-(*tert*-Butoxyoxalyl-amino)-pyrazine-2-carboxylic acid (31.7 mg, 0.12 mmol) was stirred in 20 % trifluoroacetic acid in dichloromethane (1 ml) at room temperature for 2 h. The volatiles were evaporated in vacuo and the residue co-evaporated with toluene in vacuo affording 25 mg (100 %) of the title compound as a solid.

¹H NMR (400 MHz, CDCl₃) δ 7.80 (d, 1H, J = 1.5 Hz), 8.15 (d, 1H, J = 1.5 Hz), 8.62 (s, 1H).

EXAMPLE 36



2-(Oxalyl-amino)-nicotinic acid:

To a solution of 2-aminonicotinic acid (61.4 mg, 0.45 mmol) in tetrahydrofuran (1 ml) was added imidazol-1-yl-oxo-acetic acid *tert*-butyl ester (174.2mg, 0.89 mmol) and triethylamine (62 µl, 0.45 mmol). The mixture was stirred at room temperature for 2 h. The solvent was removed in vacuo and the residue was dissolved in ethyl acetate (5.0 ml) and washed with 1% hydrochloric acid (2 x 2 ml) then water (2 x 2 ml). The

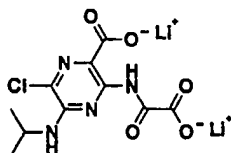
organic phase was dried (MgSO_4), filtered and the solvent evaporated in vacuo. The residue (125 mg) was purified by preparative TLC (Kieselgel 60F₂₅₄, 1mm, $\text{CH}_2\text{Cl}_2/\text{MeOH}$, 80/20) affording 7.9 mg (7 %) of 2-(*tert*-butoxyoxalyl-amino)-nicotinic acid as a solid.

5 ^1H NMR (400 MHz, CD_3OD) δ 1.80 (s, 9H), 7.40 (m, 1H), 8.50 - 8.70 (m, 2H).

2-(*tert*-Butoxyoxalyl-amino)-nicotinic acid (7.1 mg, 0.03 mmol) was stirred in 20 % trifluoroacetic acid in dichloromethane (0.5 ml) at room temperature for 2 h. The volatiles were evaporated to dryness in vacuo affording 5.6 mg (100 %) of the title compound as a solid.

10 ^1H NMR (400 MHz, CD_3OD) δ 7.40 (m, 1H), 8.50 - 8.70 (m, 2H).

EXAMPLE 37



15 6-Chloro-5-isopropylamino-3-(oxalyl-amino)-pyrazine-2-carboxylic acid, dilithium salt:

To a solution of 3-amino-6-chloro-5-isopropylamino-pyrazine-2-carboxylic acid (65.4 mg, 0.27 mmol) in tetrahydrofuran (1 ml) was added imidazol-1-yl-oxo-acetic acid *tert*-butyl ester (104.8 mg, 0.54 mmol) and triethylamine (37.4 μl , 0.27 mmol). The mixture was stirred at room temperature for 20 h followed by heating to 50 $^\circ\text{C}$ for 1.5 h. The solvent was removed in vacuo and the residue was dissolved in ethyl acetate (5.0 ml) and washed with 1 % hydrochloric acid (2 x 2 ml) then water (2 x 2 ml). The organic phase was dried (MgSO_4), filtered and the solvent evaporated in vacuo affording crude 97 mg (97 %) of 3-(*tert*-butoxyoxalyl-amino)-6-chloro-5-isopropylamino-pyrazine-2-carboxylic acid as an oil.

25 ^1H NMR (400 MHz, CDCl_3) δ 1.50 (d, 6H), 1.80 (s, 9H), 4.10 (s, 3H), 4.40 (m, 1H).

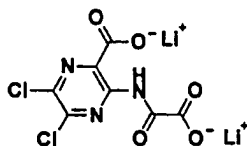
3-(*tert*-Butoxyoxalyl-amino)-6-chloro-5-isopropylamino-pyrazine-2-carboxylic acid (30 mg, 0.1 mmol) was dissolved in tetrahydrofuran (1 ml) and 1.0 N lithium hydroxide (1

ml, 1 mmol) was added at room temperature. The reaction mixture was stirred for 3 days at room temperature. After removing the solvent *in vacuo*, the residue was dissolved in ethyl acetate (20 ml) and washed with water (4 x 3.0 ml). The organic phase was dried (Na₂SO₄), filtered and the solvent evaporation *in vacuo* affording 21 mg (82 %) of the title compound as a solid.

¹H NMR (400 MHz, CD₃OD) δ 1.42 (d, 6H), 4.50 (m, 1H).

MS *m/z* 228 (M-74).

EXAMPLE 38



5,6-Dichloro-3-(oxalyl-amino)-pyrazine-2-carboxylic acid, dilithium salt:

To a solution of 3-amino-5,6-dichloro-pyrazine-2-carboxylic acid (57 mg, 0.26 mmol) in tetrahydrofuran (0.5 ml) was added imidazol-1-yl-oxo-acetic acid *tert*-butyl ester (100.6 mg, 0.513 mmol) and triethylamine (35.8 μl, 0.26 mmol). The mixture was stirred at room temperature for 20 h followed by heating at 40 °C for 4 h. The solvent was removed *in vacuo* and the residue was dissolved in ethyl acetate (5.0 ml) and washed with 1 % hydrochloric acid (2 x 2 ml) then water (2 x 2 ml). The organic phase was dried (MgSO₄), filtered and the solvent evaporated *in vacuo*. The residual oil was purified by preparative tlc (Kieselgel 60F₂₅₄, 1 mm, hexane: ethyl acetate, 1:1) affording 23.6 mg (26 %) of 3-(*tert*-butoxyoxalyl-amino)-5,6-dichloro-pyrazine-2-carboxylic acid as an oil.

¹H NMR (400 MHz, CDCl₃) δ 1.58 (s, 9H), 1.80 (s, 9H), 3.90 (s, 3H).

To a solution of 3-(*tert*-Butoxyoxalyl-amino)-5,6-dichloro-pyrazine-2-carboxylic acid (23 mg, 0.07 mmol) in tetrahydrofuran (0.5 ml) was added a 1.0 N aqueous solution of lithium hydroxide (0.5 ml) and the resulting mixture was stirred for 3 days. After removing the solvent *in vacuo*, the residue was dissolved in ethyl acetate (20 ml) and washed with water (4 x 3.0 ml). The organic phase was dried (Na₂SO₄), filtered and

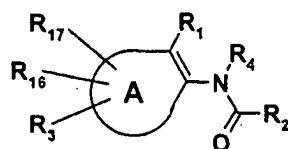
the solvent evaporation in vacuo affording 14 mg (80 %) of the title compound as a solid.

MS m/z 290.3 (M-74).

CLAIMS

1. A compound of Formula 1

5

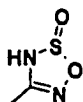
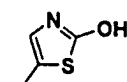
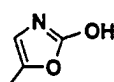
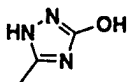
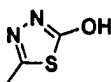
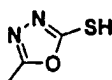
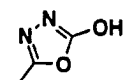
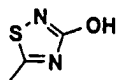
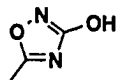
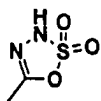
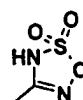
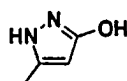
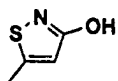
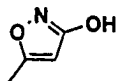
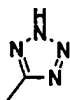


Formula 1

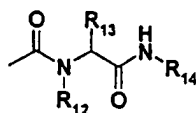
10 wherein

A is together with the double bond in Formula 1 phenyl, biphenyl, indenyl, fluorenyl, fluorenyl-9-one, naphthyl, pyridyl, pyridazinyl, pyrimidyl or pyrazinyl;

15 R_1 is hydrogen, COR_5 , OR_6 , CF_3 , nitro, cyano, SO_3H , $\text{SO}_2\text{NR}_7\text{R}_8$, $\text{PO}(\text{OH})_2$, $\text{CH}_2\text{PO}(\text{OH})_2$, $\text{CHFPO}(\text{OH})_2$, $\text{CF}_2\text{PO}(\text{OH})_2$, $\text{C}(\text{=NH})\text{NH}_2$, NR_7R_8 or selected from the following 5-membered heterocycles:



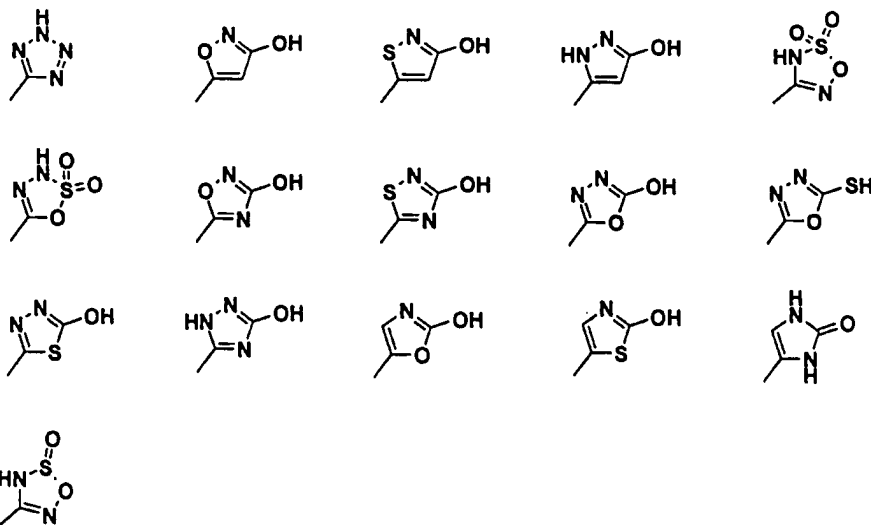
or R_1 is



5

wherein R_{12} , R_{13} , and R_{14} are independently hydrogen, C_1 - C_6 alkyl, aryl, aryl C_1 - C_6 alkyl and the alkyl and aryl groups are optionally substituted;

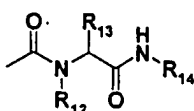
- 10 R_2 is COR_5 , OR_6 , CF_3 , nitro, cyano, SO_3H , $SO_2NR_7R_8$, $PO(OH)_2$, $CH_2PO(OH)_2$, $CHFPO(OH)_2$, $CF_2PO(OH)_2$, $C(=NH)NH_2$, NR_7R_8 , or selected from the following 5-membered heterocycles:



15

- R_3 , R_{16} and R_{17} are independently hydrogen, halo, nitro, cyano, trihalomethyl, C_1 - C_6 alkyl, aryl, aryl C_1 - C_6 alkyl, hydroxy, carboxy, carboxy C_1 - C_6 alkyl, C_1 - C_6 alkyloxycarbonyl, aryloxycarbonyl, aryl C_1 - C_6 alkyloxycarbonyl, C_1 - C_6 alkyloxy, C_1 - C_6 alkyloxy C_1 - C_6 alkyl, aryloxy, aryl C_1 - C_6 alkyloxy, aryl C_1 - C_6 alkyloxy C_1 - C_6 alkyl, thio, C_1 - C_6 alkylthio, C_1 - C_6 alkylthio C_1 - C_6 alkyl, arylthio, aryl C_1 - C_6 alkylthio, aryl C_1 - C_6 alkylthio C_1 - C_6 alkyl, NR_7R_8 , C_1 - C_6 alkylamino C_1 - C_6 alkyl, aryl C_1 - C_6 alkylamino C_1 - C_6 alkyl, di(aryl C_1 - C_6 alkyl)amino C_1 - C_6 alkyl, C_1 - C_6 alkylcarbonyl, C_1 - C_6 alkylcarbonyl- C_1 - C_6 alkyl, aryl C_1 -
- 20

C₆alkylcarbonyl, arylC₁-C₆alkylcarbonylC₁-C₆alkyl, C₁-C₆alkylcarboxy, C₁-C₆alkylcarboxyC₁-C₆alkyl, arylcarboxy, arylcarboxyC₁-C₆alkyl, arylC₁-C₆alkylcarboxy, arylC₁-C₆alkylcarboxyC₁-C₆alkyl, C₁-C₆alkyl-carbonylamino, C₁-C₆alkylcarbonylaminoC₁-C₆alkyl, -carbonylNR₇C₁-C₆alkylCOR₁₁, arylC₁-C₆alkylcarbonylamino, arylC₁-C₆alkylcarbonyl-aminoC₁-C₆alkyl, CONR₇R₈, or C₁-C₆alkylCONR₇R₈ wherein the alkyl and aryl groups are optionally substituted and R₁₁ is NR₇R₈, or C₁-C₆alkylNR₇R₈; or R₃ is



wherein R₁₂, R₁₃, and R₁₄ are independently hydrogen, C₁-C₆alkyl, aryl, arylC₁-C₆alkyl and the alkyl and aryl groups are optionally substituted;

R₄ is hydrogen, hydroxy, C₁-C₆alkyl, aryl, arylC₁-C₆alkyl, NR₇R₈, C₁-C₆alkyloxy; wherein the alkyl and aryl groups are optionally substituted;

R₅ is hydroxy, C₁-C₆alkyl, aryl, arylC₁-C₆alkyl, C₁-C₆alkyloxy, C₁-C₆alkyl-oxyC₁-

C₆alkyloxy, aryloxy, arylC₁-C₆alkyloxy, CF₃, NR₇R₈; wherein the alkyl and aryl groups are optionally substituted;

R₆ is hydrogen, C₁-C₆alkyl, aryl, arylC₁-C₆alkyl; wherein the alkyl and aryl groups are optionally substituted;

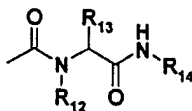
R₇ and R₈ are independently selected from hydrogen, C₁-C₆alkyl, aryl, arylC₁-C₆alkyl, C₁-C₆alkylcarbonyl, arylcarbonyl, arylC₁-C₆alkylcarbonyl, C₁-C₆alkylcarboxy or arylC₁-C₆alkylcarboxy wherein the alkyl and aryl groups are optionally substituted; or

R₇ and R₈ are taken together with the nitrogen to which they are attached forming a saturated, partially saturated or aromatic cyclic, bicyclic or tricyclic ring system containing from 3 to 14 carbon atoms and from 0 to 3 additional heteroatoms selected from nitrogen, oxygen or sulfur, the ring system can optionally be substituted with at

least one C₁-C₆alkyl, aryl, arylC₁-C₆alkyl, hydroxy, oxo, C₁-C₆alkyloxy, arylC₁-C₆alkyloxy, C₁-C₆alkyloxyC₁-C₆alkyl, NR₉R₁₀ or C₁-C₆alkylaminoC₁-C₆alkyl, wherein R₉ and R₁₀ are independently selected from hydrogen, C₁-C₆alkyl, aryl, arylC₁-C₆alkyl, C₁-C₆alkylcarbonyl, arylcarbonyl, arylC₁-C₆alkylcarbonyl, C₁-C₆alkylcarboxy or arylC₁-C₆alkylcarboxy; wherein the alkyl and aryl groups are optionally substituted; or
5 R₇ and R₈ are independently a saturated or partial saturated cyclic 5, 6 or 7 membered amine, imide or lactam;
or a salt thereof with a pharmaceutically acceptable acid or base, or any optical isomer or mixture of optical isomers, including a racemic mixture, or any tautomeric
10 forms, provided that when A is phenyl, R₁ and R₂ is COOH and R₄ is hydrogen, then R₃, R₁₆ and R₁₇ cannot be hydrogen at the same time.

2. A compound according to claim 1 wherein A is phenyl.
- 15 3. A compound according to claim 1 wherein A is naphthyl.
4. A compound according to claim 1 wherein A is indenyl.
5. A compound according to claim 1 wherein A is fluorenyl.
- 20 6. A compound according to claim 1 wherein A is fluorenyl-9-one.
7. A compound according to claim 1 wherein A is biphenyl.
- 25 8. A compound according to claim 1 wherein A is pyridyl.
9. A compound according to claim 1 wherein A is pyrimidyl.
10. A compound according to claim 1 wherein A is pyrazinyl.
- 30 11. A compound according to claim 1 wherein A is pyridazinyl.

12. A compound according to claim 2 to 11 wherein R_3 is



5

wherein R_{12} , R_{13} , and R_{14} are defined as above.

13. A compound according to claim 2 to 11 wherein R_1 and R_2 are COR_5 and R_4 is
 10 hydrogen; wherein R_5 is defined as above, provided that when A is phenyl, R_1 and R_2 is $COOH$ and R_4 is hydrogen, then R_3 , R_{16} and R_{17} cannot be hydrogen at the same time.

14. A compound according to claim 2 to 11 wherein R_1 is 5-tetrazolyl and R_2 is
 15 COR_5 , wherein R_5 is defined as above.

15. A compound according to claim 2 to 11 wherein R_1 and R_2 are $COOH$ and R_4 is
 hydrogen, provided that when A is phenyl, R_1 and R_2 is $COOH$ and R_4 is hydrogen,
 then R_3 , R_{16} and R_{17} cannot be hydrogen at the same time.

20

16. A compound selected from the following:

- 2-Methyl-6-(oxalyl-amino)benzoic acid;
- 2-(Oxalyl-amino)-5-iodo-benzoic acid;
- 25 2-(Aminooxalyl-amino)benzoic acid;
- 2-(Oxalyl-amino)-4-fluoro-benzoic acid;
- 2-(Oxalyl-amino)-4-(3-hydroxy-pyridin-2-ylsulfanyl)benzoic acid;
- 2-(Oxalyl-amino)-4-(1-imidazol-2-ylsulfanyl)benzoic acid;
- 2-(Oxalyl-amino)-4-(benzooxazol-2-ylsulfanyl)benzoic acid;
- 30 2-(Oxalyl-amino)-4-(4-phenyl-thiazol-2-ylsulfanyl)benzoic acid;
- 2-(Oxalyl-amino)-4-(3-ethoxycarbonylphenoxy)benzoic acid;

- 2-(Oxalyl-amino)-4-benzylamino-benzoic acid;
2-(Oxalyl-amino)-4-(cyclopropylmethyl-amino)benzoic acid;
2-(Oxalyl-amino)-4-(3-methyl-butylamino)benzoic acid;
2-(Hydrazinooxalyl-amino)benzoic acid;
5 3-(Oxalyl-amino)naphthalene-2-carboxylic acid;
2-((5-Thioxo-4,5-dihydro[1,3,4]oxadiazole-2-carbonyl)-amino)benzoic acid;
N-(2-(3,5-Dichloro-phenoxy)phenyl)oxalamic acid;
2-(Oxalyl-amino)-3-hydroxy-benzoic acid;
2-(Oxalyl-amino)-5-hydroxy-benzoic acid;
10 2-(Oxalyl-amino)-4-nitro-benzoic acid;
2-(Oxalyl-amino)-3,5-diiodo-benzoic acid;
2-(Oxalyl-amino)-3,5-dimethyl-benzoic acid;
2-(Oxalyl-amino)-5-methyl-benzoic acid;
2-(Oxalyl-amino)-4,5-dimethoxy-benzoic acid;
15 3-(Oxalyl-amino)-phthalic acid;
2-(Oxalyl-amino)-4-(2-hydroxy-cyclohexylamino)benzoic acid;
2-(Oxalyl-amino)-4-((2-hydroxy-ethyl)-methyl-amino)benzoic acid;
2-(Oxalyl-amino)-4-(2,2,6,6-tetramethyl-piperidin-4-ylamino)benzoic acid;
2-(2-Hydroxy-cyclohexylamino)-6-(oxalyl-amino)benzoic acid;
20 2-[(2-Hydroxy-ethyl)-methyl-amino]-6-(oxalyl-amino)benzoic acid;
2-(Oxalyl-amino)-6-(2,2,6,6-tetramethyl-piperidin-4-ylamino)benzoic acid;
2-(Oxalyl-amino)-4-(pyridin-3-ylamino)benzoic acid;
2-(Oxalyl-amino)-4-(3-hydroxy-piperidin-1-yl)benzoic acid;
2-(Oxalyl-amino)-4-(1-methyl-1H-imidazol-2-ylsulfanyl)benzoic acid;
25 2-(Oxalyl-amino)-6-(pyridin-3-ylamino)benzoic acid;
2-(3-Hydroxy-piperidin-1-yl)-6-(oxalyl-amino)benzoic acid;
2-(1-Methyl-1H-imidazol-2-ylsulfanyl)-6-(oxalyl-amino)benzoic acid;
2-(Oxalyl-amino)-4-(2-morpholin-4-yl-ethylamino)benzoic acid;
2-(Oxalyl-amino)-4-(benzenesulfonyl-(2-morpholin-4-yl-ethyl)-amino)benzoic acid;
30 2-(Oxalyl-amino)-4-[acetyl-(2-morpholin-4-yl-ethyl)-amino]benzoic acid;
2-(2-Morpholin-4-yl-ethylamino)-6-(oxalyl-amino)benzoic acid;

- 2-(Benzenesulfonyl-(2-morpholin-4-yl-ethyl)-amino)-6-(oxalyl-amino)benzoic acid;
2-(Acetyl-(2-morpholin-4-yl-ethyl)-amino)-6-(oxalyl-amino)benzoic acid;
2-(Oxalyl-amino)-4-cyclopentylamino-benzoic acid;
2-(Oxalyl-amino)-4-(benzenesulfonyl-cyclopentyl-amino)benzoic acid;
5 2-(Oxalyl-amino)-4-(acetyl-cyclopentyl-amino)benzoic acid;
2-Cyclopentylamino-6-(oxalyl-amino)benzoic acid;
2-(Benzenesulfonyl-cyclopentyl-amino)-6-(oxalyl-amino)benzoic acid;
2-(Acetyl-cyclopentyl-amino)-6-(oxalyl-amino)benzoic acid;
4-(Oxalyl-amino)-biphenyl-3-carboxylic acid;
10 3-(Oxalyl-amino)-isonicotinic acid;
5-(Oxalyl-amino)-2,6-dioxo-1,2,3,6-tetrahydro-pyrimidine-4-carboxylic acid;
2-(Oxalyl-amino)-nicotinic acid;
6-Chloro-5-isopropylamino-3-(oxalyl-amino)-pyrazine-2-carboxylic acid;
5,6-Dichloro-3-(oxalyl-amino)-pyrazine-2-carboxylic acid;
15 3-(Oxalyl-amino)-pyridazine-4-carboxylic acid;
4-(Oxalyl-amino)-pyrimidine-5-carboxylic acid;
3-(Oxalyl-amino)-pyrazine-2-carboxylic acid;

or a pharmaceutically acceptable salt thereof.

20

17. Compounds according to any one of the preceding claims which acts as inhibitors or modulators of Protein Tyrosine Phosphatases.

- 25 18. A pharmaceutical composition comprising a compound according to any of the claim 1 to 16 or a pharmaceutical acceptable salt thereof with a pharmaceutically acceptable acid or base, or any optical isomer or mixture of optical isomers, including a racemic mixture, or any tautomeric forms together with one or more pharmaceutically acceptable carriers or diluents.

30

19. A pharmaceutical composition suitable for treating type I diabetes, type II diabetes, impaired glucose tolerance, insulin resistance or obesity comprising a

compound according to any of the claims 1 to 16 or a pharmaceutical acceptable salt thereof with a pharmaceutically acceptable acid or base, or any optical isomer or mixture of optical isomers, including a racemic mixture, or any tautomeric forms together with one or more pharmaceutically acceptable carriers or diluents.

5

20. A pharmaceutical composition suitable for treating immune dysfunctions including autoimmunity, diseases with dysfunctions of the coagulation system, allergic diseases including asthma, osteoporosis, proliferative disorders including cancer and psoriasis, diseases with decreased or increased synthesis or effects of growth hormone, diseases with decreased or increased synthesis of hormones or cytokines that
10 regulate the release of/or response to growth hormone, diseases of the brain including Alzheimer's disease and schizophrenia, and infectious diseases comprising a compound according to any of the claims 1 to 16 or a pharmaceutical acceptable salt thereof with a pharmaceutically acceptable acid or base, or any optical isomer or
15 mixture of optical isomers, including a racemic mixture, or any tautomeric forms together with one or more pharmaceutically acceptable carriers or diluents.

20

21. The pharmaceutical composition according to claim 18, 19 or 20 in the form of an oral dosage unit or parenteral dosage unit.

25

22. A pharmaceutical composition according to claim 18, 19 or 20 wherein said compound is administered as a dose in a range from about 0.05 to 1000 mg, preferably from about 0.1 to 500 mg and especially in the range from 50 to 200 mg per day.

30

23. A compound according to any one of the claims 1 to 16 or a pharmaceutically acceptable salt thereof with a pharmaceutically acceptable acid or base, or any optical isomer or mixture of optical isomers, including a racemic mixture, or any tautomeric forms for therapeutical use.

24. A compound according to any one of the claims 1 to 16 or a pharmaceutically acceptable salt thereof with a pharmaceutically acceptable acid or base, or any optical isomer or mixture of optical isomers, including a racemic mixture, or any tautomeric forms for therapeutical use in the treatment or preventing of type I
5 diabetes, type II diabetes, impaired glucose tolerance, insulin resistance or obesity.

25. A compound according to any one of the claims 1 to 16 or a pharmaceutically acceptable salt thereof with a pharmaceutically acceptable acid or base, or any optical isomer or mixture of optical isomers, including a racemic mixture,
10 or any tautomeric forms for therapeutical use in the treatment or preventing of immune dysfunctions including autoimmunity, diseases with dysfunctions of the coagulation system, allergic diseases including asthma, osteoporosis, proliferative disorders including cancer and psoriasis, diseases with decreased or increased synthesis or effects of growth hormone, diseases with decreased or increased
15 synthesis of hormones or cytokines that regulate the release of/or response to growth hormone, diseases of the brain including Alzheimer's disease and schizophrenia, and infectious diseases.

26. The use of a compound according to any one of the claims 1 to 16 or a
20 pharmaceutically acceptable salt thereof with a pharmaceutically acceptable acid or base, or any optical isomer or mixture of optical isomers, including a racemic mixture, or any tautomeric forms as a medicament.

27. The use of a compound according to any of the claims 1 to 16 for preparing
25 a medicament.

28. The use of a compound according to any one of the claims 1 to 16 or a pharmaceutically acceptable salt thereof with a pharmaceutically acceptable acid or base, or any optical isomer or mixture of optical isomers, including a racemic mixture,
30 or any tautomeric forms for the preparation of a medicament suitable for the

treatment or preventing of type I diabetes, type II diabetes, impaired glucose tolerance, insulin resistance or obesity.

29. The use of a compound according to any one of the claims 1 to 16 or a
5 pharmaceutically acceptable salt thereof with a pharmaceutically acceptable acid or base, or any optical isomer or mixture of optical isomers, including a racemic mixture, or any tautomeric forms for the preparation of a medicament suitable for the treatment or preventing of immune dysfunctions including autoimmunity, diseases with dysfunctions of the coagulation system, allergic diseases including asthma,
10 osteoporosis, proliferative disorders including cancer and psoriasis, diseases with decreased or increased synthesis or effects of growth hormone, diseases with decreased or increased synthesis of hormones or cytokines that regulate the release of/or response to growth hormone, diseases of the brain including Alzheimer's disease and schizophrenia, and infectious diseases.

15

30. A method of treating type I diabetes, type II diabetes, impaired glucose tolerance, insulin resistance or obesity comprising administering to a subject in need thereof an effective amount of a compound according to any of the claims 1 to 16 to said subject.

20

31. A method of treating immune dysfunctions including autoimmunity, diseases with dysfunctions of the coagulation system, allergic diseases including asthma, osteoporosis, proliferative disorders including cancer and psoriasis, diseases with decreased or increased synthesis or effects of growth hormone, diseases with
25 decreased or increased synthesis of hormones or cytokines that regulate the release of/or response to growth hormone, diseases of the brain including Alzheimer's disease and schizophrenia, and infectious diseases comprising administering to a subject in need thereof an effective amount of a compound according to any of the claims 1 to 16 to said subject.

30

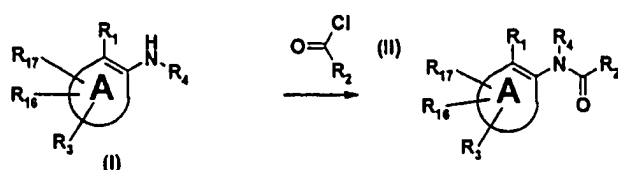
32. A process for the manufacture of a medicament, particular to be used in the treatment or prevention of type I diabetes, type II diabetes, impaired glucose tolerance, insulin resistance or obesity which process comprising bringing a compound according to any of the claims 1 to 16 or a pharmaceutically acceptable salt thereof into a galenic dosage form.

33. A process for the manufacture of a medicament, particular to be used in the treatment or prevention of immune dysfunctions including autoimmunity, diseases with dysfunctions of the coagulation system, allergic diseases including asthma, osteoporosis, proliferative disorders including cancer and psoriasis, diseases with decreased or increased synthesis or effects of growth hormone, diseases with decreased or increased synthesis of hormones or cytokines that regulate the release of/or response to growth hormone, diseases of the brain including Alzheimer's disease and schizophrenia, and infectious diseases which process comprising bringing a compound according to any of the claims 1 to 16 or a pharmaceutically acceptable salt thereof into a galenic dosage form.

34. Any novel feature or combination of features as described herein.

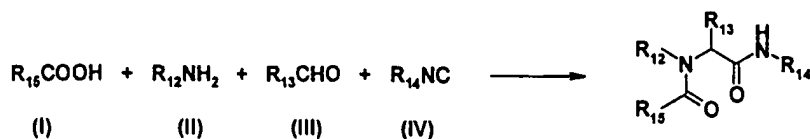
35. A method of preparing a compound of formula 1, characterized in

a)

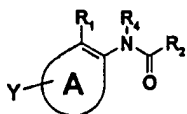


allowing an amino substituted compound of formula (I) to react with an acid chloride of formula (II), wherein A, R₁, R₂, R₃, R₄, R₁₆ and R₁₇ are defined as above, or

b)



allowing a carboxylic acid (I), a primary amine (II) and an aldehyde (III) to react with a
 isocyanide (IV) wherein R_{12} , R_{13} , R_{14} , and R_{15} are independently selected from the
 group consisting of hydrogen, C_1 - C_6 alkyl, aryl, aryl C_1 - C_6 alkyl as defined above and
 the alkyl and aryl groups are optionally substituted as defined above; or
 R_{12} , R_{13} , R_{14} , and R_{15} are independently selected from



wherein Y indicates attachment point for R_{12} , R_{13} , R_{14} , and R_{15} and A, R_1 , R_2 and R_4
 are defined as above, or

c)

the above described four component Ugi reaction (method b)) is carried out by at-
 taching any one of the components to a solid support whereby the synthesis is ac-
 complished in a combinatorial chemistry fashion.

36. Compounds according to claim 1 to 16 which acts as ligands, inhibitors or
 modulators of molecules with pTyr recognition units including proteins that contain
 SH2 domains.

1
INTERNATIONAL SEARCH REPORT

International application No.
PCT/DK 99/00122

A. CLASSIFICATION OF SUBJECT MATTER

IPC6: C07C 235/84, C07D 213/55, C07D 239/28, C07D 241/24, A61K 31/17, A61K 31/44,
A61K 31/50, A61K 31/505
According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC6: C07C, C07D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE,DK,FI,NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	DE 2413966 A (AMERICAN HOME PRODUCTS CORP.), 26 Sept 1974 (26.09.74) --	1
X	US 3987192 A (JOHN B. WRIGHT), 19 October 1976 (19.10.76) --	1
X	US 4169153 A (JOHN B. WRIGHT), 25 Sept 1979 (25.09.79) --	1
X	US 4457872 A (KIYOSHI MURASE ET AL), 3 July 1984 (03.07.84) --	1

☒ Further documents are listed in the continuation of Box C. ☒ See patent family annex.

* Special categories of cited documents:	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A" document defining the general state of the art which is not considered to be of particular relevance	"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"E" earlier document but published on or after the international filing date	"Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"&" document member of the same patent family
"O" document referring to an oral disclosure, use, exhibition or other means	
"P" document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

28 June 1999

Date of mailing of the international search report

13 -07- 1999

Name and mailing address of the ISA/
Swedish Patent Office
Box 5055, S-102 42 STOCKHOLM
Facsimile No. +46 8 666 02 86

Authorized officer
Göran Karlsson/Els
Telephone No. +46 8 782 25 00

INTERNATIONAL SEARCH REPORT

International application No.

PCT/DK 99/00122

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 3966965 A (JOHN H. SELLSTEDT ET AL), 29 June 1976 (29.06.76) --	1
X	GB 1493338 A (AMERICAN HOME PRODUCTS CORPORATION), 30 November 1977 (30.11.77) --	1
X	EP 0030436 A1 (YAMANOUCHI PHARMACEUTICAL CO. LTD.), 17 June 1981 (17.06.81) --	1
X	GB 1449934 A (AMERICAN HOME PRODUCTS CORPORATION), 15 Sept 1976 (15.09.76) --	1
X	Chemical Abstracts, Volume 93, No 10, 8 Sept 1980 (08.09.80), (Columbus, Ohio, USA), Bezuglyi, P.A. et al, "N-Heteryloxamines 1-adamantylammonium having diuretic activity", page 391, THE ABSTRACT No 101484m, Obraztsy, Tovarnye Znaki 1980 (7), 116-117 --	1
X	Chemical Abstracts, Volume 93, No 24, 15 December 1980 (15.12.80), (Columbus, Ohio, USA), Bezuglyi, P.A. et al, "Tert-Butylammonium N-heteryloxamates possessing diuretic activity", page 318, THE ABSTRACT No 225644h, Obraztsy, Tovarnye Znaki 1980 (22), 144-145 -- -----	1

INTERNATIONAL SEARCH REPORT

International application No.
PCT/DK99/00122

Box I Observations where certain claims were found unsearchable (Continuation of Item 1 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.: 26-27, 30-31
because they relate to subject matter not required to be searched by this Authority, namely:
**A method for treatment of the human or animal body by therapy,
see rule 39.1.**
2. ☒ Claims Nos.: 1-25, 28-29, 32-36
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
see next page

Box II Observations where unity of invention is lacking (Continuation of Item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims: it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/DK99/00122

The present claims are drafted in such a complex and broad way that no meaningful search is possible to perform, cf. Article 6. The multitude of variables and their permutations and combinations result in a claimed subject matter that is so broad in scope that it is rendered virtually incomprehensible.

A preliminary search of a part of claim 1 has shown that a great number of known compounds are included in this claim. Many of these compounds have the effects given in claims 20 and 29.

The claimed subject matter lacks a significant structural element qualifying as a special technical feature that clearly defines a contribution over the prior art. Due to the lack of such a feature and the complexity of the claims, the present claims state a great number of separate inventions. No invitation to pay additional fees has been made as this would have meant a unreasonable number of fees.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.

01/06/99

PCT/DK 99/00122

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
DE 2413966 A	26/09/74	AR 207563 A	15/10/76
		AU 6575374 A	21/08/75
		BE 812743 A	23/09/74
		CA 1034118 A	04/07/78
		CH 612914 A	31/08/79
		EG 11281 A	31/05/77
		FI 61688 B,C	31/05/82
		FR 2222097 A,B	18/10/74
		GB 1449934 A	15/09/76
		GB 1449935 A	15/09/76
		GB 1449936 A	15/09/76
		GB 1493337 A	30/11/77
		GB 1493338 A	30/11/77
		IE 39336 B	13/09/78
		IE 39337 B	13/09/78
		IE 39338 B	13/09/78
		JP 1248338 C	16/01/85
		JP 49135929 A	27/12/74
		JP 59020664 B	15/05/84
		NL 7403843 A	25/09/74
		PH 14870 A	08/01/82
		SE 411898 B,C	11/02/80
		US 3966965 A	29/06/76
		US 4036837 A	19/07/77
		US 4044144 A	23/08/77
		US 4054657 A	18/10/77
		US 4054661 A	18/10/77
		US 4054666 A	18/10/77
		US 4069343 A	17/01/78
		US 4154961 A	15/05/79
		US 4160100 A	03/07/79
		US 4191840 A	04/03/80
		ZA 7400635 A	24/09/75
US 3987192 A	19/10/76	NONE	
US 4169153 A	25/09/79	CA 1066714 A	20/11/79
		DE 2525226 A,C	18/12/75
		FR 2273523 A,B	02/01/76
		GB 1514775 A	21/06/78
		GB 1514776 A	21/06/78
		JP 1136363 C	28/02/83
		JP 51006941 A	20/01/76
		JP 57026595 B	05/06/82
		US 4091011 A	23/05/78
US 4457872 A	03/07/84	AT 7385 E,T	15/05/84
		CA 1181070 A	15/01/85
		EP 0030436 A,B	17/06/81
		JP 57064652 A	19/04/82
		JP 57064651 A	19/04/82
		JP 57075953 A	12/05/82
		JP 1516752 C	07/09/89
		JP 56077247 A	25/06/81
		JP 63062509 B	02/12/88

INTERNATIONAL SEARCH REPORT
Information on patent family members

01/06/99

International application No.
PCT/DK 99/00122

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 3966965 A	29/06/76	GB 1493337 A	30/11/77
		GB 1493338 A	30/11/77
		JP 51141834 A	07/12/76
		US 4036837 A	19/07/77
		US 4044144 A	23/08/77
		US 4054657 A	18/10/77
		US 4054661 A	18/10/77
		US 4054666 A	18/10/77
		US 4069343 A	17/01/78
		US 4154961 A	15/05/79
		US 4160100 A	03/07/79
		US 4191840 A	04/03/80
		AR 207563 A	15/10/76
		AU 6575374 A	21/08/75
		BE 812743 A	23/09/74
		CA 1034118 A	04/07/78
		CH 612914 A	31/08/79
		DE 2413966 A	26/09/74
		EG 11281 A	31/05/77
		FI 61688 B,C	31/05/82
		FR 2222097 A,B	18/10/74
		GB 1449934 A	15/09/76
		GB 1449935 A	15/09/76
		GB 1449936 A	15/09/76
		IE 39336 B	13/09/78
		IE 39337 B	13/09/78
		IE 39338 B	13/09/78
		JP 1248338 C	16/01/85
		JP 49135929 A	27/12/74
		JP 59020664 B	15/05/84
		NL 7403843 A	25/09/74
		PH 14870 A	08/01/82
		SE 411898 B,C	11/02/80
		ZA 7400635 A	24/09/75

INTERNATIONAL SEARCH REPORT
Information on patent family members

01/06/99

International application No.

PCT/DK 99/00122

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
GB 1493338 A	30/11/77	GB 1449934 A	15/09/76
		GB 1449935 A	15/09/76
		GB 1449936 A	15/09/76
		GB 1493337 A	30/11/77
		AR 207563 A	15/10/76
		AU 6575374 A	21/08/75
		BE 812743 A	23/09/74
		CA 1034118 A	04/07/78
		CH 612914 A	31/08/79
		DE 2413966 A	26/09/74
		EG 11281 A	31/05/77
		FI 61688 B,C	31/05/82
		FR 2222097 A,B	18/10/74
		IE 39336 B	13/09/78
		IE 39337 B	13/09/78
		IE 39338 B	13/09/78
		JP 1248338 C	16/01/85
		JP 49135929 A	27/12/74
		JP 59020664 B	15/05/84
		NL 7403843 A	25/09/74
		PH 14870 A	08/01/82
		SE 411898 B,C	11/02/80
		US 3966965 A	29/06/76
		US 4036837 A	19/07/77
		US 4044144 A	23/08/77
		US 4054657 A	18/10/77
		US 4054661 A	18/10/77
		US 4054666 A	18/10/77
		US 4069343 A	17/01/78
		US 4154961 A	15/05/79
		US 4160100 A	03/07/79
		US 4191840 A	04/03/80
		ZA 7400635 A	24/09/75
		JP 51141834 A	07/12/76
EP 0030436 A1	17/06/81	AT 7385 E,T	15/05/84
		CA 1181070 A	15/01/85
		JP 57064652 A	19/04/82
		US 4457872 A	03/07/84
		JP 57064651 A	19/04/82
		JP 57075953 A	12/05/82
		JP 1516752 C	07/09/89
		JP 56077247 A	25/06/81
		JP 63062509 B	02/12/88

INTERNATIONAL SEARCH REPORT
Information on patent family members

01/06/99

International application No.

PCT/DK 99/00122

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
GB 1449934 A	15/09/76	GB 1449935 A	15/09/76
		GB 1449936 A	15/09/76
		GB 1493337 A	30/11/77
		GB 1493338 A	30/11/77
		AR 207563 A	15/10/76
		AU 6575374 A	21/08/75
		BE 812743 A	23/09/74
		CA 1034118 A	04/07/78
		CH 612914 A	31/08/79
		DE 2413966 A	26/09/74
		EG 11281 A	31/05/77
		FI 61688 B,C	31/05/82
		FR 2222097 A,B	18/10/74
		IE 39336 B	13/09/78
		IE 39337 B	13/09/78
		IE 39338 B	13/09/78
		JP 1248338 C	16/01/85
		JP 49135929 A	27/12/74
		JP 59020664 B	15/05/84
		NL 7403843 A	25/09/74
		PH 14870 A	08/01/82
		SE 411898 B,C	11/02/80
		US 3966965 A	29/06/76
		US 4036837 A	19/07/77
		US 4044144 A	23/08/77
		US 4054657 A	18/10/77
		US 4054661 A	18/10/77
		US 4054666 A	18/10/77
		US 4069343 A	17/01/78
		US 4154961 A	15/05/79
		US 4160100 A	03/07/79
		US 4191840 A	04/03/80
		ZA 7400635 A	24/09/75